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(71) Applicant: **JOHNSON AND JOHNSON CONSUMER COMPANIES, INC.** [US/US]; 199 Grandview Road, Skillman, NJ 08558 (US).

(72) Inventors: **LUKENBACH, Elvin, R.**; 160 Klinesville Road, Flemington, NJ 08822 (US). **KAMINSKI, Claudia**; 490 Shire Road, Milford, NJ 08848 (US). **PASCAL-SUISSE, Sandrine**; Residence Les Annonciades, 5 bis, rue du Mont, F-76000 Rouen (FR). **TAHAR, Maurice**; 72, rue de la Marne, F-27200 Vernon (FR).

(74) Agents: **CIAMPORCERO, Audley, A. et al.**; Johnson and Johnson, One Johnson and Johnson Plaza, New Brunswick, NJ 08933 (US).

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(54) Title: **CLEANSING COMPOSITIONS**

(57) Abstract: Cleansing compositions suitable for use in personal cleansing applications, and in particular make-up removal applications, which not only impart superior cleansing properties, but also which are relatively non-irritating and thus suitable for use by people having sensitive skin and eyes comprised of esters, liquid silicones, and a water dispersible components. Also disclosed are compositions for effectively depositing various benefit agents into and onto the skin.

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Cleansing Compositions

BACKGROUND OF THE INVENTION

5 1. Field of the Invention

This invention relates to cleansing compositions suitable for use in personal cleansing applications, and in particular make-up removal applications, which not only impart superior cleansing properties, but also which are relatively non-irritating and thus suitable for use by
10 people having sensitive skin and eyes. This invention is further related to a composition for effectively delivering and/or depositing various benefit agents into and onto the skin.

2. Description of the Prior Art

Various types of cosmetics such as make-up, e.g. lipstick, mascara, foundation, and the like, leave an oil-containing residue on the skin surface that cannot be removed easily by
15 facial cleansers containing conventional soaps. One reason is that such soaps are unable to effectively emulsify or solubilize such oils, which is why many make-up remover compositions have included an oil base as a major component. For example, European Patent No. 370856 discloses a non-foaming makeup remover system comprised of a surfactant-containing water phase that remains physically separated from a cosmetic oil-containing oil phase
20 unless shaken. Disadvantageously, such oil-containing removers also suffer from a tendency to deposit an oily residue or film on the user's skin.

Various attempts have been made to produce stable, oil-free makeup removers. For example, United States Patent No. 5,217,641 discloses an oil-free, stable, non-irritating, single-phase makeup remover comprised of 50 percent to 98 percent of cyclomethicone along with a
25 mixture of esters. However, not only is it economically disadvantageous to use such a large amount of cyclomethicones, but because of their highly volatile nature, cyclomethicones cannot be packaged easily using conventional cosmetic packaging.

Another reason that make-up cannot be removed by conventional soaps is the fact that such soaps are incapable of removing the binders in the make-up. These binders tend to
30 increase the make-up's resistance against sebum and water as well as its overall adhesiveness to the skin. In addition, various polymers, which are similarly difficult to remove, are also employed in hair cosmetics for the purpose of protecting the hair or providing the hair with body.

It would be desirable to have a stable, economically-feasible composition that could not
35 only effectively remove the residue from sebum as well as the residue from make-up and hair-

protecting agents, but also impart a pleasant, non-oily "after-feel" to the skin and hair. It would also be desirable to create such a composition having a low degree of ocular and skin irritation. It would further be desirable to create such a composition that is capable of depositing various active agents into and onto the skin.

5 SUMMARY OF THE INVENTION

In accordance with this invention, there is provided a cleaning composition comprising:

- a. a liquid silicone;
- b. a water dispersible component; and
- c. an ester.

10 Another embodiment of this invention is directed to a cleaning system comprising:

- a. a liquid silicone;
- b. a water dispersible component;
- c. an ester;
- d. water; and
- 15 e. a polymeric emulsifier and/or thickener.

Another embodiment of this invention is directed to a foaming composition comprising:

- a. a water dispersible component;
- b. an ester;
- c. water; and a
- 20 d. foaming surfactant.

Yet another embodiment of the present invention is directed to a method for making an oil-in water emulsion comprised of:

neutralizing a hydrophilic thickening agent in a hydrophilic phase comprised of a polymeric emulsifier with an effective amount of a neutralizer under conditions sufficient after
25 a lipophilic phase was combined with the hydrophilic phase.

Yet another embodiment of the present invention is directed to a method for making a water-in-oil emulsion comprised of:

neutralizing a hydrophilic thickening agent in a hydrophilic phase comprised of a polymeric emulsifier with an effective amount of a neutralizer under conditions sufficient
30 before combining a lipophilic phase with the hydrophilic phase.

Yet another embodiment of the present invention is directed to a method for depositing benefit agents into and onto the skin comprised of:

5 topically applying an effective amount of the benefit agent with a composition comprised of an optional liquid silicone, a water dispersible component, and an ester to a desired location.

Yet another embodiment of the present invention is directed to a method for depositing a benefit agent into and/or onto the skin, hair and/or nails comprising applying a composition comprising:

- 10 a. an optional liquid silicone;
- b. a water dispersible component;
- c. an ester;
- d. a polymeric emulsifier and/or thickener; and
- e. an effective amount of a benefit agent

to a desired location on a human or animal.

15 Yet another embodiment of the present invention is directed to a method for depositing a benefit agent into and/or onto the skin, hair and/or nails comprising applying a composition comprising:

- a. an optional liquid silicone;
- b. a water dispersible component
- 20 c. an ester;
- d. water;
- e. a foaming surfactant; and
- f. an effective amount of a benefit agent

to a desired location on a human or animal.

25 The compositions of this invention are capable of effectively cleansing the skin without significantly contributing to ocular irritation as well as depositing various benefit agents into and onto the skin, hair and nails.

DESCRIPTION OF THE DRAWINGS

30 The file of this patent contains at least one drawing executed in color. Copies of this patent with color drawing(s) will be provided by the Patent and Trademark Office upon request and payment of the necessary fee.

The invention will be more fully understood and further advantages will become apparent when reference is made to the following detailed description of the invention and the accompanying drawing in which:

FIG. 1 is a representation that illustrates two forearms, each of which contains three sets of three different types of makeup before a cleanser is applied thereto.

FIGS. 2 (a) through (f) are representations of the same forearm after six respective cleansers were applied to each set of makeup, respectively.

FIGS. 3 (a) and (b) are representations that illustrate the right side (FIG 3(a)) and left side (FIG. 3(b)) of a subject's face prior to treatment as viewed under a CG-395 Filter.

FIGS. 3(c) and (d) are representations that illustrate the right side (FIG 3(c)) and left side (FIG. 3(d)) of a subject's face while possessing the formulation of Example 10 as viewed under a CG-395 Filter.

FIG. 3(e) is a representation that illustrates the right side of a subject's face after the treatment of Example 10 was rinsed therefrom as viewed under a CG-395 Filter.

FIG. 3(f) is a representation that illustrates the left side of a subject's face after the treatment of Example 10 was wiped therefrom as viewed under a CG-395 Filter.

FIGS. 4 (a) and (b) are representations that illustrate the right side (FIG 4(a)) and left side (FIG. 4(b)) of a subject's face prior to treatment as viewed under a CG-395 Filter.

FIGS. 4(c) and (d) are representations that illustrate the right side (FIG 4(c)) and left side (FIG. 4(d)) of a subject's face while possessing the formulation of Example 10 as viewed under a CG-395 Filter.

FIG. 4(e) is a representation that illustrates the right side of a subject's face after the treatment of Example 10 was rinsed therefrom as viewed under a CG-395 Filter.

FIG. 4(f) is a representation that illustrates the left side of a subject's face after the treatment of Example 10 was wiped therefrom as viewed under a CG-395 Filter.

FIG. 5(a) is a graph of concentration of retinol in the formulation of Example 10 versus pixel intensity change.

FIG. 5(b) is a graph of concentration of retinol in the formulation of Example 11 versus pixel intensity change.

DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

In one embodiment of the present invention, the cleaning composition may suitably comprise, consist of, or consist essentially of, based upon the total weight of the cleaning composition, a) from about 10 percent to about 35 percent, and preferably from about 10 percent to about 20 percent of a liquid silicone; b) from about 10 percent to about 35 percent, and preferably from about 10 percent to about 20 percent of a water dispersible component; and c) from about 30 percent to about 80 percent, and preferably from 55 percent to about 65 percent of an ester.

The first component of the cleaning composition of the present invention may be either a volatile or nonvolatile liquid silicone, with the former being preferred. Examples of suitable silicones nonexclusively include the polydimethyl siloxanes and derivatives thereof such as hexamethylsiloxane, dimethicone, dimethiconol, and cyclomethicone, with cyclomethicone being preferred. Examples of suitable cyclomethicones nonexclusively include cyclotetradimethyl siloxane; cyclopentadimethyl siloxane, cyclohexadimethyl siloxane, cycloheptadimethyl siloxane, and mixtures thereof. Preferably, the silicone has a viscosity of from about 0.25 cp to about 350 cp.

The second component of the cleaning composition of the present invention is a water dispersible component, which is preferably a water soluble solvent. As used herein, the term "water dispersible component" shall mean a material that produces a uniform, clear or hazy, mixture when combined with at least a weight equivalent of water. Examples of suitable water dispersible components nonexclusively include polyethylene glycol 400, hexylene glycol, propylene glycol, polypropylene glycol-10 methylglucose ether, ethoxydiglycol, polyethylene glycol-6 caprylic/capric glyceride, ethylene glycol monobutyl ether, polyethylene glycol-8 caprylic/capric glycerides, 3-methoxy-3-methyl-1-butanol, dimethyl isosorbide, and mixtures thereof. Most preferred water dispersible components include hexylene glycol, dimethyl isosorbide, polyethylene glycol-6 caprylic/capric glyceride, and mixtures thereof.

The third component of the cleaning composition of the present invention is a lipophilic component that preferably is a liquid ester. Preferred esters for use in the composition of this invention include those liquid esters that either possess a structural means for ensuring its liquidity or are heterogeneous in nature. Examples of such structural means include the presence of "interruptions", such as: 1) chain branching; 2) olefinic unsaturation; 3) the presence of either a polyether or a monoalkoxylate in the structure; or 4) the presence of a substituent, e.g. an ethoxylated or propoxylated moiety, bonded between the acid group and the alcohol group. By "heterogeneity," it is meant that the lipophilic component is comprised of a mixture of compounds that vary in the number of carbon atoms in their respective chains.

Examples of suitable esters nonexclusively include:

- a) a branched C₅ to C₂₂ alkyl alcohol ester of an aromatic acid;
 - b) a straight-chained or branched C₅ to C₂₂ alkyl acid esters of optionally ethoxylated/propoxylated polyols having from about 3 carbon atoms to about 7 carbon atoms;
 - c) branched C₅ to C₂₂ alkyl alcohol esters of branched polyacids;
 - d) branched or straight-chained C₅ to C₂₂ alkyl acid esters of branched and/or unsaturated C₅ to C₂₂ alkyl alcohols;
 - e) branched or unsaturated C₅ to C₂₂ alkyl alcohol esters of an acid selected from the group consisting of adipic acid, succinic acid, maleic acid, sebacic acid, and mixtures thereof;
 - f) polyether interrupted fatty acid esters;
 - g) benzoic acid ester of heterogeneous alcohols having from about 8 carbon atoms to about 22 carbon atoms; and
 - h) mixtures thereof,
- with straight-chained or branched C₅ to C₂₂ alkyl acid esters of optionally ethoxylated/propoxylated polyols, benzoic acid esters of heterogeneous alcohols, and mixtures thereof being particularly preferred.

Suitable branched C₅ to C₂₂ alkyl alcohol esters of an aromatic acid include those wherein the aromatic acid is benzoic acid. Preferably, the alcohol of this ester is either branched or unsaturated, and may be either a primary alcohol or a secondary alcohol with the former being preferred. Optionally, the aromatic acid may be substituted with hydroxy or alkyl groups having from about 1 carbon atom to about 4 carbon atoms. Specific examples of these esters nonexclusively include, butyloctyl salicylate; hexyldecyl benzoate; and butyloctyl benzoate, which are all available from C.P. Hall Co. under the tradename, "HallStar;" and mixtures thereof, with a mixture of hexyldecyl benzoate and butyloctyl benzoate being particularly preferred.

Another suitable ester includes a straight-chained or branched C₅ to C₂₂ alkyl acid ester of optionally ethoxylated/propoxylated polyols, wherein the polyols contain from about 3 carbon atoms to about 7 carbon atoms. Preferably, if the polyol creates a branching point, then the acid group may be straight-chained. Suitable acids used to form these esters typically have from about 8 carbon atoms to about 22 carbon atoms, and preferably from about 8 carbon atoms to about 18 carbon atoms, and most preferably from about 8 carbon atoms to about 12 carbon atoms, and are either saturated or unsaturated, with octanoic acid,

capric acid, and mixtures thereof being preferred. Such suitable acids are either straight-chained or branched, and are preferably aliphatic. Suitable polyols used to form these esters typically have from about 3 carbon atoms to about 30 carbon atoms, and preferably from about 3 carbon atoms to about 7 carbon atoms. Examples of such suitable polyols nonexclusively include neopentyl alcohol; polyglycerol, e.g. diglycerol, triglycerol, hexaglycerol, and decaglycerol, wherein the polyglycerol may contain from about 2 to about 10 glycerol groups; glycerin; sorbitan; methyl glucose; trimethylolpropane; and mixtures thereof. Neopentyl alcohol, glycerin, trimethylolpropane, and mixtures thereof are the preferred polyols. Examples of suitable esters nonexclusively include pentaerythritol tetraoctanoate; trimethylolpropane trioctanoate; trioctanoin; pentaerythrityl tetrapelargonate; sorbitan trioleate; caprylic/capric triglyceride; neopentyl alcohol tetraoctanoate, and mixtures thereof, with caprylic/capric triglyceride; pentaerythritol tetraoctanoate; trimethylolpropane trioctanoate; and pentaerythrityl tetrapelargonate being more preferred.

Another suitable ester includes the branched C_5 to C_{22} alkyl alcohol esters of branched polyacids such as the tri-esters, tetra-esters, penta-esters, and mixtures thereof. An example of such a polyacid is citric acid. Suitable alkyl alcohols for creating these esters are optionally substituted, e.g., ethoxylated or propoxylated, contain from about 3 carbon atoms to about 22 carbon atoms, and preferably from about 3 carbon atoms to about 8 carbon atoms, and are either straight-chained or branched, with the branching being preferred. These alcohols may either be primary or secondary, and may either be saturated or unsaturated, with saturated being preferred for stability reasons. Specific examples of suitable esters nonexclusively include trioctyldodecyl citrate; triisopropylcitrate; and mixtures thereof.

Another suitable ester includes the branched or straight-chained C_5 to C_{22} alkyl acid esters of branched or unsaturated alkyl alcohols wherein the alkyl group of the alcohol has from about 1 carbon atoms to about 18 carbon atoms, and preferably from about 4 carbon atoms to about 10 carbon atoms, provided that the total number of carbon atoms in the ester is at least about 8. Suitable acids for use in making these esters typically have from about 2 carbon atoms to about 22 carbon atoms, and preferably from about 5 carbon atoms to about 10 carbon atoms. However, if the number of acid carbon atoms exceeds the number of carbon atoms in the alcohol, then the acid preferably contains from about 8 carbon atoms to about 18 carbon atoms and the alcohol preferably contains from about 1 carbon atom to about 8 carbon atoms. If the number of acid carbon atoms is less than the number of carbon atoms in the alcohol, then the acid preferably contains from about 2 carbon atoms to about 8 carbon atoms and the alcohol preferably contains from about 8 carbon atoms to about 18 carbon atoms. Preferably, either: 1) the alcohol group or the acid group has branching and/or unsaturation, i.e. both the alcohol and the acid are not straight-chained; or 2) the

ester possesses an asymmetrical alkyl distribution. By "asymmetrical alkyl distribution," it is meant that the ester is made from, for example, a short chain alcohol, i.e. having from about 1 carbon atom to about 8 carbon atoms, and a long chain acid, i.e., having greater than about 8 carbon atoms, such as, e.g. butyl stearate, or less preferably the ester is made from, a long chain alcohol, i.e. having greater than about 8 carbon atoms, and a short chain acid, i.e. having from about 1 carbon atom to about 8 carbon atoms. Examples of such suitable esters nonexclusively include tridecyl neopentanoate, isostearyl palmitate, cetyl ricinoleate, cetyl octanoate, isononyl isononanoate, butyl stearate, octyldodecyl soyate, tridecyl erucate, octyldodecyl erucate/eicosil erucate, and mixtures thereof, with cetyl octanoate, isostearyl palmitate, isononyl isononanoate, and mixtures thereof and being preferred.

Another suitable ester includes the branched or unsaturated C₅ to C₂₂ alkyl alcohol esters of an acid selected from the group consisting of adipic acid, succinic acid, maleic acid, sebacic acid, and mixtures thereof. The alcohol of these esters, which has from about 3 carbon atoms to about 18 carbon atoms, and preferably from about 3 carbon atoms to about 8 carbon atoms, is preferably branched or unsaturated. Examples of such suitable alcohol esters nonexclusively include diisopropyl adipate, dioctyl sebacate, dioctyl succinate, dioctyl maleate, diisostearyl adipate, diethyl sebacate, and mixtures thereof, with diethyl sebacate, dioctyl sebacate, and diisostearyl adipate being preferred.

Another suitable ester includes polyether interrupted fatty acid esters. Examples of such suitable esters nonexclusively include: 1) laureth-2 benzoate; 2) C₈ to C₂₂ fatty alkyl (optionally polypropylenoxy) polyethyleneoxy carboxylate esters derived from an alcohol having from about 1 carbon atom to about 22 carbon atoms, is either straight or branched, and may contain a phenyl group; and 3) mixtures thereof, with C₈ to C₂₂ fatty alkyl (optionally polypropylenoxy) polyethyleneoxy carboxylate esters being preferred. Specific examples of preferred esters nonexclusively include isopropyl propylene glycol-2-isodeceth-7 carboxylate, such as "Velsan D8P3" and other commercially available materials sold by Sandoz under the tradename, "Velsan."

Another suitable ester includes the benzoic acid esters of heterogeneous alcohols having from about 8 carbon atoms to about 22 carbon atoms, such as the ester mixtures available from Finetex under the tradename, "Finsolv" and preferably is the C₁₂ to C₁₅ alcohols benzoate available from Finetex under the tradename, "Finsolv TN."

Preferred combinations of esters include at least one, preferably at least two, and more preferably three of the following esters: a) branched C₅ to C₂₂ alkyl alcohol esters of an aromatic acid; b) branched or straight-chained C₅ to C₂₂ alkyl acid esters of branched or unsaturated alkyl alcohols; and c) straight-chained or branched C₅ to C₂₂ alkyl acid esters of optionally ethoxylated/propoxylated polyols. In a preferred embodiment, the ester

contains, based upon the total weight percent of the esters, from about 30 percent to about 80 percent of branched or straight-chained C₅ to C₂₂ alkyl acid esters of branched or unsaturated C₅ to C₂₂ alkyl alcohols; from about 10 percent to about 50 percent of branched C₅ to C₂₂ alkyl alcohol esters of an aromatic acid; and from about 10 percent to about 50 percent of straight-chained or branched C₅ to C₂₂ alkyl acid esters of optionally ethoxylated/propoxylated polyols. In a more preferred embodiment, the ester contains, based upon the total weight percent of the esters, from about 15 percent to about 50 percent isononyl isononanoate, from about 15 percent to about 50 percent isostearyl palmitate, from about 15 percent to about 50 percent cetyl octanoate, and from about 15 percent to about 50 percent pentaerthritol tetraoctanoate.

Another embodiment of the present invention is directed to a cleaning system comprising, consisting, or consisting essentially of, based upon the total weight of the cleaning system, a) at least 5 percent and preferably at least about 10 percent of the cleaning composition; b) from about 70 percent to about 98 percent, and preferably from about 80 percent to about 90 percent of water; c) from about 0.1 percent to 5 percent, e.g. from about 0.5 percent to about 1.5 percent of a polymeric emulsifier, a thickener, or mixture thereof; optionally d) from about 0.1 percent to about 5 percent, and preferably from about 1 percent to about 3 percent of a cleaning enhancer; optionally e) from about 2 percent to about 20 percent, and preferably from about 5 percent to about 15 percent of a foaming surfactant; and optionally f) from about 0.001 percent to about 20 percent of a benefit agent. In one embodiment, the cleaning system may comprise, based upon the total weight of the cleansing system, from about 0.1 to about 5 percent, and preferably from about 0.5 percent to 1.5 percent of a polymeric emulsifier and/or from about 0.01 percent to about 2 percent, and preferably from about 0.01 percent to about 0.5 percent of a thickener. More preferably, the cleaning system contains, based upon the total weight of the cleaning system, from about 10 percent to about 30 percent of the cleaning composition.

The cleaning system may be in the form of an oil-in-water emulsion, a water-in-oil emulsion, or a dispersion.

In addition to the cleansing composition, the cleaning system is further comprised of polymeric emulsifiers and/or thickeners. As used herein, the term "polymeric emulsifier" shall mean those compounds capable of emulsifying cleaning systems whereby the polymeric emulsifiers have a molecular weight of at least about 5000, and preferably are block copolymers having a hydrophilic portion and a hydrophobic portion. When used at amounts effective for emulsifying the cleansing system, the polymeric emulsifiers surprisingly do not cause significant eye sting, i.e., when the emulsifier-containing composition was used by 80 consumers in the eye area, no more than about 5% of such users expressed discomfort

around the eye area. Examples of suitable polymeric emulsifiers nonexclusively include polyethylene glycol-30 dipolyhydroxystearate available from Uniqema under the tradename, 'Arlacel P-135;" dimethicone copolyol, which is available from Goldschmidt Chemical Corporation under the tradename, "Abil EM 90"; substituted acrylates such as those available
5 from The Goodrich Corporation under the tradename, "Pemulen"; and mixtures thereof, with polyethylene glycol-30 dipolyhydroxystearate being preferred.

Examples of suitable hydrophilic thickeners nonexclusively include carbomers available from B.F. Goodrich under the tradename, "Carbopol ETD 2020", acrylate copolymers, hydroxyethylcellulose modified with cetyl ether groups available from Hercules
10 under the tradename, "Natrosol Plus", polyvinylmethyl ether/maleic anhydride (PVM/MA) decadiene crosspolymer available from International Specialty Products under the tradename, "Stabileze QM," and copolymers and mixtures thereof, with carbomers being preferred. Examples of suitable acrylate copolymers nonexclusively include acrylate copolymers available from Rohm & Haas under the tradename, "Aculyn 33,"
15 acrylates/aminoacrylates copolymer available from National Starch & Chemical Company under the tradename, "Structure Plus," acrylates/steareth-20 itaconate copolymer available from National Starch & Chemical Company under the tradename, "Structure 2001," acrylates/ceteth-20 itaconate copolymer available from National Starch & Chemical Company under the tradename, "Structure 3001," acrylates/steareth-20 methacrylate
20 copolymer available from Rohm & Haas under the tradename, "Aculyn 22," and copolymers and mixtures thereof.

The cleaning system of the present invention may also optionally contain a cleaning enhancer in the form of a nonionic emulsifier and/or a non-foaming surfactant. Examples of suitable nonionic emulsifiers include isocetheth-20, oleth-2, mixture of PEG-40 hydrogenated
25 castor oil and trideceth-9 available from Dragoco Inc. under the tradename, "Dragoco Solubilizer 2/014160," Poloxamer 184, laureth-4, sorbitan trioleate, polyoxyethylene-(2) oleyl ether, sorbitan stearate, cetearyl glucoside, glyceryl oleate, trideceth-9, polyethylene glycol-40 hydrogenated castor oil, and mixtures thereof.

Examples of suitable non-foaming surfactants include non-foaming nonionic
30 surfactants such as sucrose esters, e.g., sucrose cocoate, sucrose stearate and mixtures thereof, with sucrose cocoate being preferred. By "essentially non-foaming," it is meant that the surfactant, when used with the composition of the present invention, has a column height of less than about 20 mm as determined by the Ross-Miles Foam Generation Test. See 18 (l.) Oil & Soap 99 – 102 (1941) ["Ross-Miles Test"], which is incorporated by reference
35 herein. The cleansing composition and the cleansing system may either be rinseable with water or may be wiped-off. Preferably, the essentially, non-foaming surfactants are used in

embodiments wherein the cleansing system or the cleansing composition is rinseable with water. For example, a preferred combination of hydrophilic components include, based upon the total weight percent of the cleansing system, from about 0.1 percent to about 5.0 percent of hexylene glycol, from about 0.5 percent to about 3.0 percent of sucrose cocoate
5 non foaming surfactant, and from about 0.5 percent to about 3.0 percent of polyoxyethylene-6 caprylic/capric triglyceride. An example of a suitable cleaning enhancer include a mixture of sorbitan stearate and sucrose cocoate available from Uniqema under the tradename, "Arlatone 2121."

Preferably, the cleaning system contains, based upon the total weight of the
10 cleansing system, no more than about 6%, and preferably 5%, of the cleaning enhancers for cream formulations and no more than about 2%, and preferably no more than 1% of the cleaning enhancers in thin lotion/milk formulations.

The cleansing system and cleansing composition may also optionally contain a foaming surfactant. The foaming surfactant may be non-ionic, cationic, amphoteric, or
15 anionic; nonionic surfactants are preferred. By "foaming," it is meant that the surfactant, when used with the composition of the present invention, has a column height of foam greater than about 20 mm as determined by the Ross-Miles Test. As used herein, the term "amphoteric" shall mean: 1) molecules that contain both acidic and basic sites such as, for example, an amino acid containing both amino (basic) and acid (e.g., carboxylic acid, acidic)
20 functional groups; or 2) zwitterionic molecules which possess both positive and negative charges within the same molecule. The charges of the latter may be either dependent on or independent of the pH of the composition. Examples of zwitterionic materials include, but are not limited to, alkyl betaines and amidoalkyl betaines. Examples of suitable and preferred surfactants may be found in International Patent Application Number WO97/01196, which is
25 incorporated by reference in its entirety herein.

The cleansing system may further contain one or more benefit agents or pharmaceutically-acceptable salts thereof. As used herein, the term "benefit agent" includes any active ingredient that is to be delivered into and/or onto the skin, hair or nail at a desired location, such as a cosmetic agent or a pharmaceutical agent. By "cosmetic agent," it is
30 meant any ingredient that is appropriate for cosmetically treating, providing nutrients to, and/or conditioning the hair, nail, and/or skin via topical application. By "pharmaceutical agent," it is mean any drug that is either hydrophobic or hydrophilic in nature and appropriate for topical use. As used herein "medicament agents" include those agents capable of promoting recovery from injury and illness.

35 The benefit agents useful herein may be categorized by their therapeutic benefit or their postulated mode of action. However, it is to be understood that the benefit agents

useful herein may, in some circumstances, provide more than one therapeutic benefit or operate via greater than one mode of action. Therefore, the particular classifications provided herein are made for the sake of convenience and are not intended to limit the benefit agents to the particular application(s) listed. In addition, the compounds, which are
5 identified below as being suitable for use as benefit agents, may be used in an amount over and above the amount that they may be used for other purposes in the cleansing composition/cleansing system.

Examples of suitable benefit agents include, but are not limited to, depigmentation agents; reflectants; detangling/wet combing agents; film forming polymers; humectants;
10 amino acids and their derivatives; antimicrobial agents; allergy inhibitors; anti-acne agents; anti-aging agents; anti-wrinkling agents, antiseptics; analgesics; antitussives; antipruritics; local anesthetics; anti-hair loss agents; hair growth promoting agents; hair growth inhibitor agents, antihistamines such as *Mandragora Vernalis*, *Tanacetum Parthenium* and the like; anti-infectives such as *Acacia Catechu*, *Aloe Barbadensis*, *Convallaria Majalis*, *Echinacea*,
15 *Eucalyptus*, *Mentha Piperita*, *Rosa Canina*, *Sassafras Albidum*, and the like; inflammation inhibitors; anti-emetics; anticholinergics; vasoconstrictors; vasodilators; wound healing promoters; peptides, polypeptides and proteins; deodorants and anti-perspirants; medicament agents; skin emollients and skin moisturizers; skin firming agents, hair conditioners; hair softeners; hair moisturizers; vitamins; tanning agents; skin lightening
20 agents; antifungals such as *Centaurea Cyanus*, *Kalmia Latifolia* and antifungals for foot preparations; depilating agents; shaving preparations; external analgesics; perfumes; counterirritants; hemorrhoidals; insecticides; poison ivy products; poison oak products; burn products; anti-diaper rash agents; prickly heat agents; make-up preparations; vitamins; amino acids and their derivatives; herbal extracts; retinoids; flavenoids; sensates; anti-
25 oxidants; skin conditioners; hair lighteners; chelating agents; cell turnover enhancers; coloring agents; pigments; sunscreens, those active ingredients disclosed in United States Patent No. 6,063,397, which is incorporated herein by reference, anti-edema agents, collagen enhancers, and mixtures thereof.

Examples of suitable anti-edema agents nonexclusively include bisabolol natural,
30 synthetic bisabolol, and mixtures thereof.

Examples of suitable vasoconstrictors nonexclusively include horse chestnut extract, prickly ash, and mixtures thereof.

Examples of suitable anti-inflammatory agents nonexclusively include benoxaprofen, centella asiatica, bisabolol, feverfew (whole), feverfew (parthenolide free), green tea extract,
35 green tea concentrate, hydrogen peroxide, lycopene including "Lyc-o-Pen" available from LycoRed Natural Products Industries, Ltd., oat oil, chamomile, and mixtures thereof.

Examples of collagen enhancers nonexclusively include vitamin A, vitamin C, and mixtures thereof.

Examples of suitable skin firming agent nonexclusively include dimethylaminoethanol ("DMAE").

5 Examples of suitable antipruritics and skin protectants nonexclusively include oatmeal, betaglucan, feverfew, soy and derivatives thereof, bicarbonate of soda, colloidal oatmeal, surfactant based colloidal oatmeal cleanser, *Anagallis Arvensis*, *Oenothera Biennis*, *Verbena Officinalis*, and the like. These antipruritics may be used in an amount, based upon the total weight of the cleansing composition, from about 0.01 percent to about
10 40 percent, and preferably from about 1 percent to about 5 percent.

As used herein, colloidal oatmeal means the powder resulting from the grinding and further processing of whole oat grain meeting United States Standards for Number 1 or Number 2 oats. The colloidal oatmeal has a particle size distribution as follows: not more than 3 percent of the total particles exceed 150 micrometers in size and not more than 20
15 percent of the total particles exceed 75 micrometers in size. Examples of suitable colloidal oatmeals include, but are not limited to, "Tech-O" available from the Beacon Corporation and colloidal oatmeals available from Quaker.

Examples of suitable reflectants nonexclusively include mica, alumina, calcium silicate, glycol dioleate, glycol distearate, silica, sodium magnesium fluorosilicate, and
20 mixtures thereof.

Suitable detangling/wet combing agents nonexclusively include polyquaternium-10, hydroxypropyltrimonium guar, dioleoylamidoethyl hydroxyethylmonium methosulfate, di-
(soyoylethyl) hydroxyethylmonium methosulfate, hydroxyethyl behenamidopropyl dimonium
chloride, olealkonium chloride, polyquaternium-47, stearalkonium chloride, tricetylmonium
25 chloride, and mixtures thereof.

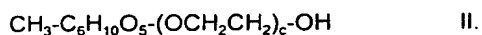
Suitable film forming polymers include those that, upon drying, produce a substantially continuous coating or film on the hair, skin, or nails. Nonexclusive examples of suitable film forming polymers include acrylamidopropyl trimonium chloride/acrylamide copolymer; corn
starch/ acrylamide/ sodium acrylate copolymer; polyquaternium-10; polyquaternium-47;
30 polyvinylmethylether/maleic anhydride copolymer; styrene/acrylates copolymers; and mixtures thereof.

Commercially available humectants which are capable of providing moisturization and conditioning properties to the cleansing composition are suitable for use in the present invention. The humectant is preferably present in an amount of from about 0 percent to about
35 10 percent, more preferably from about 0.5 percent to about 5 percent, and most preferably from about 0.5 percent to about 3 percent, based on the overall weight of the composition. Examples of suitable humectants nonexclusively include: 1) water soluble liquid polyols

selected from the group comprising glycerine, propylene glycol, hexylene glycol, butylene glycol, pentylene glycol, dipropylene glycol, and mixtures thereof; 2) polyalkylene glycol of the formula I.:



5 wherein R'' is an alkylene group having from about 2 to about 4 carbon atoms and b is an integer of from about 1 to about 10, such as PEG 4; 3) polyethylene glycol ether of methyl glucose of formula II.:



 wherein c is an integer from about 5 to about 25;

10 4) urea; 5) fructose; 6) glucose; 7) honey; 8) lactic acid; 9) maltose; 10) sodium glucuronate; and 11) mixtures thereof, with glycerine being the preferred humectant.

 Suitable amino acid agents include amino acids derived from the hydrolysis of various proteins as well as the salts, esters, and acyl derivatives thereof. Examples of such amino acid agents nonexclusively include amphoteric amino acids such as alkylamido
15 alkylamines, i.e. stearyl acetyl glutamate, capryloyl silk amino acid, capryloyl collagen amino acids; capryloyl keratin amino acids; capryloyl pea amino acids; cocodimonium hydroxypropyl silk amino acids; corn gluten amino acids; cysteine; glutamic acid; glycine; hair keratin amino acids; amino acids such as aspartic acid, threonine, serine, glutamic acid, proline, glycine, alanine, cystine, valine, methionine, isoleucine, leucine, tyrosine,
20 phenylalanine, cysteic acid, lysine, histidine, arginine, cysteine, tryptophan, citrulline; lysine; silk amino acids, wheat amino acids; and mixtures thereof

 Suitable proteins include those polymers that have a long chain, i.e. at least about 10 carbon atoms, and a high molecular weight, i.e. at least about 1000, and are formed by self-condensation of amino acids. Nonexclusive examples of such proteins include collagen,
25 deoxyribonuclease, iodized corn protein; milk protein; protease; serum protein; silk; sweet almond protein; wheat germ protein; wheat protein; alpha and beta helix of keratin proteins; hair proteins, such as intermediate filament proteins, high-sulfur proteins, ultrahigh-sulfur proteins, intermediate filament-associated proteins, high-tyrosine proteins, high-glycine tyrosine proteins, trichohyalin, and mixtures thereof.

30 Examples of suitable vitamins nonexclusively include vitamin B complex; including thiamine, nicotinic acid, biotin, pantothenic acid, choline, riboflavin, vitamin B6, vitamin B12, pyridoxine, inositol, carnitine; vitamins A,C,D,E,K and their derivatives such as vitamin A palmitate and pro-vitamins, e.g. (i.e. panthenol (pro vitamin B5) and panthenol triacetate) and mixtures thereof.

Examples of suitable antibacterial agents nonexclusively include bacitracin, erythromycin, neomycin, tetracycline, chlortetracycline, benzethonium chloride, phenol, and mixtures thereof.

5 Examples of suitable skin emollients and skin moisturizers nonexclusively include mineral oil, lanolin, vegetable oils, isostearyl isostearate, glyceryl laurate, methyl gluceth-10, methyl gluceth-20 chitosan, and mixtures thereof.

Examples of suitable hair conditioners nonexclusively include quaternized compounds such as behenamidopropyl PG-dimonium chloride, tricetylmonium chloride, dihydrogenated tallowamidoethyl hydroxyethylmonium methosulfate, and mixtures thereof as
10 well as lipophilic compounds like cetyl alcohol, stearyl alcohol, hydrogenated polydecene, and mixtures thereof.

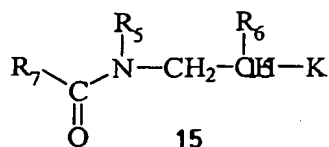
An example of a suitable hair softener nonexclusively includes silicone compounds, such as those that are either non-volatile or volatile and those that are water soluble or water insoluble. Examples of suitable silicones include organo-substituted polysiloxanes, which
15 are either linear or cyclic polymers of monomeric silicone/oxygen monomers and which nonexclusively include cetyl dimethicone; cetyl triethylammonium dimethicone copolyol phthalate; cyclomethicone; dimethicone copolyol; dimethicone copolyol lactate; hydrolyzed soy protein/dimethicone copolyol acetate; silicone quaternium 13; stearammonium dimethicone copolyol phthalate; stearamidopropyl dimethicone; and mixtures thereof.

20 Examples of suitable hair moisturizers nonexclusively include panthenyl ethyl ether, phytantriol, and mixtures thereof.

Examples of sunscreen agents nonexclusively include benzophenones, borrelone, butyl paba, cinnamidopropyl trimethyl ammonium chloride, disodium distyrylbiphenyl disulfonate, paba, potassium methoxycinnamate, butyl methoxydibenzoylmethane, octyl
25 methoxycinnamate, oxybenzone, octocrylene, octyl salicylate, phenylbenzimidazole sulfonic acid, ethyl hydroxypropyl aminobenzoate, menthyl anthranilate, aminobenzoic acid, cinoxate, diethanolamine methoxycinnamate, glyceryl aminobenzoate, titanium dioxide, zinc oxide, oxybenzone, Padimate O, red petrolatum, and mixtures thereof.

30 An example of a suitable tanning agent nonexclusively includes dihydroxyacetone. Examples of skin lightening agents nonexclusively include hydroquinone, catechol and its derivatives, ascorbic acid and its derivatives, and mixtures thereof.

Examples of suitable insecticides (including insect repellents, anti-scabies and anti-lice treatments) nonexclusively include permethrin, pyrethrin, piperonyl butoxide, imidacloprid, N,N-diethyl toluamide, which refers to the material containing predominantly the
35 *meta* isomer, i.e., N,N-diethyl-*m*-toluamide, which is also known as DEET; compounds of the formula III.



III.

wherein

- 5 R_5 is a branched or unbranched alkyl group having about 1 to about 6 carbon atoms;
- R_6 is H, methyl or ethyl;
- R_7 is a branched or unbranched alkyl or alkoxy group having from about 1 to about 8 carbon atoms; and
- 10 K is a -CN or a -COOR₈ group, wherein
- R_8 is a branched or unbranched alkyl group having from about 1 to about 6 carbon atoms,

- natural or synthetic pyrethroids, whereby the natural pyrethroids are contained in pyrethrum, the extract of the ground flowers of *Chrysanthemum cinerariaefolium* or *C. coccineum*; and
- 15 mixtures thereof. Within the structure of Formula III. are ethyl 3-(N-butylacetamido)propionate, wherein R_7 is a CH₃ group, R_5 is an n-butyl group, R_6 is H, K is COOR₈ and R_8 is ethyl, which is available commercially from Merck KGaA of Darmstadt, Germany under the name, "Insect Repellent 3535."

- An example of an anti fungal for foot preparations nonexclusively includes tolnaftate.
- 20 Examples of suitable depilating agents nonexclusively include calcium thioglycolate, magnesium thioglycolate, potassium thioglycolate, strontium thioglycolate, and mixtures thereof.

- Examples of suitable external analgesics and local anesthetics nonexclusively include benzocaine, dibucaine, benzyl alcohol, camphor, capsaicin, capsicum, capsicum
- 25 oleoresin, juniper tar, menthol, methyl nicotinate, methyl salicylate, phenol, resorcinol, turpentine oil, and mixtures thereof.

 Examples of suitable antiperspirants and deodorants nonexclusively include aluminium chlorohydrates, aluminium zirconium chlorohydrates, and mixtures thereof.

- Examples of suitable counterirritants nonexclusively include camphor, menthol,
- 30 methyl salicylate, peppermint and clove oils, ichtammol, and mixtures thereof.

 An example of a suitable inflammation inhibitor nonexclusively includes hydrocortisone, *Fragaria Vesca*, *Matricaria Chamomilla*, and *Salvia Officinalis*.

 Examples of suitable hemorrhoidal products nonexclusively include the anesthetics such as benzocaine, pramoxine hydrochloride, and mixtures thereof; antiseptics such as

benzethonium chloride; astringents such as zinc oxide, bismuth subgallate, balsam Peru, and mixtures thereof; skin protectants such as cod liver oil, vegetable oil, and mixtures thereof.

Most preferred benefit agents nonexclusively include DMAE, soy and derivatives thereof, colloidal oatmeal, sulfonated shale oil, olive leaf, elubiol, 6-(1-piperidiny)-2,4-pyrimidinediamine-3-oxide, finasteride, ketoconazole, salicylic acid, zinc pyrithione, coal tar, benzoyl peroxide, selenium sulfide, hydrocortisone, sulfur, menthol, pramoxine hydrochloride, tricetylmonium chloride, polyquaternium 10, panthenol, panthenol triacetate, vitamin A and derivatives thereof, vitamin B and derivatives thereof, vitamin C and derivatives thereof, vitamin D and derivatives thereof, vitamin E and derivatives thereof, vitamin K and derivatives thereof, keratin, lysine, arginine, hydrolyzed wheat proteins, hydrolyzed silk proteins, octyl methoxycinnamate, oxybenzone, minoxidil, titanium dioxide, zinc dioxide; retinol, erythromycin, tretinoin, and mixtures thereof.

One preferred type of benefit agent includes those therapeutic components that are effective in the treatment of dandruff, seborrheic dermatitis, and psoriasis as well as the symptoms associated therewith. Examples of such suitable benefit agents nonexclusively include zinc pyrithione, anthralin, shale oil and derivatives thereof such as sulfonated shale oil, selenium sulfide, sulfur; salicylic acid; coal tar; povidone-iodine, imidazoles such as ketoconazole, dichlorophenyl imidazolodioxalan, which is commercially available from Janssen Pharmaceutica, N.V., under the tradename, "Elubiol", clotrimazole, itraconazole, miconazole, climbazole, tioconazole, sulconazole, butoconazole, fluconazole, miconazole nitrate and any possible stereo isomers and derivatives thereof; piroctone olamine (Octopirox); selenium sulfide; ciclopirox olamine; anti-psoriasis agents such as vitamin D analogs, e.g. calcipotriol, calcitriol, and tacalcitol; vitamin A analogs such as esters of vitamin A, e.g. vitamin A palmitate, retinoids, retinols, and retinoic acid; corticosteroids such as hydrocortisone, clobetasone, butyrate, clobetasol propionate and mixtures thereof.

The amount of benefit agent to be combined with the cleansing composition or the emulsion may vary depending upon, for example, the ability of the benefit agent to penetrate through the skin, hair or nail, the specific benefit agent chosen, the particular benefit desired, the sensitivity of the user to the benefit agent, the health condition, age, and skin, hair, and/or nail condition of the user, and the like. In sum, the benefit agent is used in a "safe and effective amount," which is an amount that is high enough to deliver a desired skin, hair or nail benefit or to modify a certain condition to be treated, but is low enough to avoid serious side effects, at a reasonable risk to benefit ratio within the scope of sound medical judgment. Unless otherwise expressed herein, typically the benefit agent is present in the cleansing system in an amount, based upon the total weight of the system, from about 0.01

percent to about 20 percent, and preferably from about 0.01 percent to about 5.0 percent, and more preferably from about 0.01 percent to about 2.0 percent.

Optionally, commercially available detergent thickeners that are capable of imparting the appropriate viscosity to conditioning shampoo compositions are suitable for use in this invention. If used, the detergent thickeners should be present in the shampoo compositions in an amount sufficient to raise the Brookfield viscosity of the composition to a value of between about 500 to about 10,000 centipoise. Examples of suitable detergent thickeners nonexclusively include: mono or diesters of polyethylene glycol of formula IV.



wherein z is an integer from about 3 to about 200;

fatty acids containing from about 16 to about 22 carbon atoms; fatty acid esters of ethoxylated polyols; ethoxylated derivatives of mono and diesters of fatty acids and glycerine; hydroxyalkyl cellulose; alkyl cellulose; hydroxyalkyl alkyl cellulose; and mixtures thereof. More specifically, suitable detergent thickeners nonexclusively include behenalkonium chloride; cetyl alcohol, quaternium-46, hydroxyethyl cellulose, cocodimonium chloride, polyquaternium-6, polyquaternium-7, quaternium-18, PEG-18 glycerol oleate/cocoate, a mixture of acrylates/stearate-50 acrylate copolymer, laureth-3 and propylene glycol, which is commercially available from Goldschmidt under the tradename "Antil 208," a mixture of cocamidopropylbetaine and glyceryl laurate which is commercially available from Goldschmidt under the tradename, "Antil HS60," a mixture of propylene glycol, PEG 55, and propylene glycol oleate, which is commercially available from Goldschmidt under the tradename, "Antil 414 liquid," and mixtures thereof. Preferred detergent thickeners include polyethylene glycol ester, and more preferably PEG-150 distearate which is available from the Stepan Company of Northfield, Illinois or from Comiel, S.p.A. of Bologna, Italy under the tradename, "PEG 6000 DS".

The above described cleansing composition and cleaning system may be prepared by combining the desired components in a suitable container and mixing them under ambient conditions in any conventional mixing means well known in the art, such as a mechanically stirred propeller, paddle, and the like.

In another preferred embodiment of the cleaning system of the present invention wherein a polymeric emulsifier such as, for example, polyethylene glycol-30 dipolyhydroxystearate (hereinafter "PEG 30") or dimethicone copolyol, are used and water is used as the vehicle, an oil-in-water emulsion may be produced. Although both the PEG 30 and dimethicone copolyol are marketed for use in formulating water-in-oil compositions, we have unexpectedly found that oil-in-water emulsions may be created due to the unique processing steps and conditions employed herein. More specifically, we found that when a

thickening agent, preferably a hydrophilic thickening agent, is neutralized in the hydrophilic phase of the present invention comprising a polymeric emulsifier prior to adding the lipophilic phase of the present invention thereto, the resulting emulsion is in the form of a water-in-oil emulsion. Conversely, when a thickening agent, preferably a hydrophilic thickening agent, is neutralized in the hydrophilic phase of the present invention comprising a polymeric emulsifier after the lipophilic phase of the present invention is added to the hydrophilic phase, the resulting emulsion is unexpectedly in the form of a oil-in-water emulsion.

Cleansing systems of the present invention that are emulsions may contain, based upon the total weight of the emulsion, from about 0.01 percent to about 2 percent, and preferably from about 0.01 percent to about 0.5 percent of hydrophilic thickeners. Suitable neutralizers include any known bases, such as sodium hydroxide, or acids, such as lactic acid, that are capable of neutralizing the hydrophilic thickening agent, in either the hydrophilic phase (if a water-in-oil emulsion is desired) or a mixture of both the hydrophilic phase and the lipophilic phase (if an oil-in-water emulsion is desired) of the present invention to a pH of about 5 to about 7 under ambient temperature. In one embodiment, hydrophilic thickeners including acrylates/aminoacrylates copolymer, acrylates/steareth-20 itaconate copolymer, acrylates/ceteth-20 itaconate copolymer, are preferably neutralized with an acid, such as lactic acid. Hydrophilic thickeners including carbomers, modified hydroxyethylcellulose, polyvinylacetate/maleic anhydride (PVA/MA) decadiene crosspolymer, and acrylates/steareth-20 methacrylate copolymer, are preferably neutralized with a base, such as sodium hydroxide (20%).

In one embodiment, the hydrophilic phase may be comprised of one or more of the following components: water, thickener, cleansing enhancer, nonfoaming surfactant, and water dispersible component, and the lipophilic phase may be comprised of one or more of the following components: silicone, ester, and polymeric emulsifier.

We have also surprisingly found that the cleansing system of the present invention possesses good aesthetic properties without causing any significant ocular discomfort to the user. It is well-known in the art that most emulsifiers having a relatively low molecular weight are irritating regardless of their hydrophilic lipophilic balance ("HLB") value. However, we have surprisingly found that when the cleansing system of the present invention is produced using the particular polymeric emulsifiers and/ or thickeners set forth herein, the resulting cleanser is gentle and possesses a low degree of ocular and skin irritation.

Another embodiment of this invention is directed to a foaming composition comprising, based upon the foaming composition, from about 0.1 percent to about 30 percent, e.g. from about 0.1 percent to about 5 percent of a water dispersible component; from about 0.1 percent to about 30 percent, e.g. from about 0.1 percent to about 5 percent of an ester; from

about 1.0 percent to about 98 percent, e.g. from about 30 percent to about 98 percent or from about 45 percent to about 90 percent of water; and from about 2.0 percent to about 20 percent, e.g. from about 5.0 percent to about 15 percent of a foaming surfactant.

Optionally, the foaming composition may also be comprised of one or more of the following components, based upon the total weight of the foaming composition: a) from about 0.1 percent to 5 percent, e.g. from about 0.5 percent to about 1.5 percent of a polymeric emulsifier, a thickener, or mixture thereof; b) from about 0.1 percent to about 5 percent, e.g. from about 1 percent to about 3 percent of a cleaning enhancer; c) from about 0.001 percent to about 20 percent of a benefit agent; and d) from about 0.1 percent to about 30 percent, e.g. from about 0.1 percent to about 5 percent of a liquid silicone.

Another embodiment of the present invention is directed to a method for depositing a benefit agent onto the skin, hair and/or nails comprised of applying either the above-described cleansing system or cleansing composition with an effective amount of a benefit agent to a desired location on a human or animal. While the frequency and amount of the benefit agent-containing cleaning system to be applied will depend upon, for example, the type and amount of benefit agent available, the intended usage of the final composition, i.e. therapeutic versus maintenance regimen, the amount and type of detergent present, and the sensitivity of the individual user to the composition/emulsion, typically the benefit agent-containing cleaning system of the present invention should be topically applied to affected body parts at regular intervals, and preferably from about 2 to about 14 times per week. More preferably, the composition/emulsion is applied more frequently during the initial stages of treatment, e.g. from about 5 to about 7 times per week until the desired effect is achieved, then less frequently when maintenance is desired, e.g. from about 2 to about 5 times per week.

We have unexpectedly found that the above-described cleansing composition and cleansing system are capable of efficiently mediating the deposition and permeation of various benefit agents, such as antidandruff agents, onto and into the skin following topical administration thereto. More specifically, we have surprisingly found that when benefit agents are combined with either the cleansing composition or the cleaning system of the present invention, the amount of benefit agents deposited onto and/or into the skin, hair, and/or nails is about 50% greater than the amount of benefit agents deposited onto and/or into the skin, hair, and/or nails after application of known, commercial benefit agent-containing cleansers.

An alternative preferred embodiment of the present invention is directed to a method for treating hair loss, such as hair loss resulting from alopecia, comprising topically applying the above-described cleaning system and the hair loss benefit agent to a desired location on an animal or human, wherein the benefit agent is comprised of an effective amount of a hair

loss treatment agent such as minoxidil or mixture thereof. As used herein, "hair loss treatment agents" shall include agents capable of growing hair and/or agents capable of preventing the loss of hair. By "effective amount," it is meant an amount effective for treating hair loss and preferably may range from, based upon the total weight of the cleansing system, from about 0.001 percent to about 20 percent, and preferably from about 1 percent to about 5 percent.

Examples of benefit agents suitable for treating hair loss include, but are not limited to potassium channel openers or peripheral vasodilators such as minoxidil, diazoxide, and compounds such as N'-cyano-N-(tert-pentyl)-N'-3-pyridinyl-guanidine ("P-1075") as disclosed in United States Patent No.: 5,244,664, which is incorporated herein by reference; vitamins, such as vitamin E and vitamin C, and derivatives thereof such as vitamin E acetate and vitamin C palmitate; hormones, such as erythropoietin, prostaglandins, such as prostaglandin E₁ and prostaglandin F₂-alpha; fatty acids, such as oleic acid; diuretics such as spironolactone; heat shock proteins ("HSP"), such as HSP 27 and HSP 72; calcium channel blockers, such as verapamil HCL, nifedipine, and diltiazemamiloride; immunosuppressant drugs, such as cyclosporin and Fk-506; 5 alpha-reductase inhibitors such as finasteride; growth factors such as, EGF, IGF and FGF; transforming growth factor beta; tumor necrosis factor; non-steroidal anti-inflammatory agents such as benoxaprofen; retinoids such as tretinoin; cytokines, such as IL-6, IL-1 alpha, and IL-1 beta; cell adhesion molecules such as ICAM; glucocorticoids such as betametasone; botanical extracts such as aloe, clove, ginseng, rehmannia, swertia, sweet orange, zanthoxylum, Serenoa repens (saw palmetto), Hypoxis rooperi, stinging nettle, pumpkin seeds, and rye pollen; other botanical extracts including sandalwood, red beet root, chrysanthemum, rosemary, burdock root and other hair growth promoter activators which are disclosed in DE 4330597 which is incorporated by reference in its entirety herein; homeopathic agents such as Kalium Phosphoricum D2, Azadirachta indica D2, and Joborandi D1; genes for cytokines, growth factors, and male-patterned baldness; antifungals such as ketoconazole and elubiol; antibiotics such as streptomycin; proteins inhibitors such as cycloheximide; acetazolamide; benoxaprofen; cortisone; diltiazem; hexachlorobenzene; hydantoin; nifedipine; penicillamine; phenothiazines; pinacidil; psoralens, verapamil; zidovudine; alpha-glucosylated rutin having at least one of the following rutins: quercetin, isoquercitrin, hespeddin, naringin, and methylhesperidin, and flavonoids and transglycosidated derivatives thereof which are all disclosed in JP 7002677, which is incorporated by reference in its entirety herein; and mixtures thereof.

Preferred hair loss treatment agents include minoxidil, 6-(1-piperdiny)-2,4-pyrimidinediamine-3-oxide, N'-cyano-N-(tert-pentyl)-N'-3-pyridinyl-guanidine, finasteride, retinoids and derivatives thereof, ketoconazole, elubiol or mixtures thereof.

Another embodiment of the present invention is directed to a method for inhibiting hair growth comprising topically applying the above-described composition/system combined with a benefit agent to a desired area on an animal or human for inhibiting hair growth, wherein the benefit agent is comprised of an effective amount of a hair growth inhibiting agent. In a preferred embodiment, the cleaning system contains, based upon the total weight of the cleaning system, from about 0.001 percent to about 20 percent, and preferably from about 0.01 percent to about 5 percent hair growth inhibiting agent.

Examples of benefit agents suitable for use in inhibiting hair growth include: serine proteases such as trypsin; vitamins such as alpha-tocophenol (vitamin E) and derivatives thereof such as tocophenol acetate and tocophenol palmitate; antineoplastic agents, such as doxorubicin, cyclophosphamide, chlormethine, methotrexate, fluorouracil, vincristine, daunorubicin, bleomycin and hydroxycarbamide; anticoagulants, such as heparin, heparinoids, coumaerins, detran and indandiones; antithyroid drugs, such as iodine, thiouracils and carbimazole; lithium and lithium carbonate; interferons, such as interferon alpha, interferon alpha-2a and interferon alpha-2b; retinoids, such as retinol (vitamin A), isotretinoin; glucocorticoids such as betamethasone, and dexamethosone; antihyperlipidaemic drugs, such as triparanol and clofibrate; thallium; mercury; albendazole; allopurinol; amiodarone; amphetamines; androgens; bromocriptine; butyrophenones; carbamazepine; cholestyramine; cimetidine; clofibrate; danazol; desipramine; dixyrazine; ethambutol; etionamide; fluoxetine; gentamicin, gold salts; hydantoins; ibuprofen; imipramine; immunoglobulins; indandiones; indomethacin; intraconazole; levadopa; maprotiline; methysergide; metoprolol; metyrapone; nadolol; nicotinic acid; potassium thiocyanate; propranolol; pyridostimine; salicylates; sulfasalazine; terfenadine; thiamphenicol; thiouracils; trimethadione; tropanolol; valproic acid; and mixtures thereof.

Preferred hair growth inhibitory agents include serine proteases, retinol, isotretinoin, betamethoisone, alpha-tocophenol and derivatives thereof, or mixtures thereof.

Another preferred embodiment of the present invention is directed to a method for treating acne and for reducing the signs of aging, i.e. wrinkles, fine lines, and other manifestations of photodamage, comprising topically applying the above-described cleaning system and the relevant benefit agent to the skin of an animal or human at a desired area, wherein the benefit agent is comprised of an effective amount of an anti-acne agent or an anti-aging agent, respectively.

Examples of suitable anti-aging agents include, but are not limited to inorganic sunscreens such as titanium dioxide and zinc oxide; organic sunscreens such as octyl-methoxy cinnamates and derivatives thereof; retinoids; vitamins such as vitamin E, vitamin A, vitamin C, vitamin B, and derivatives thereof such as vitamin E acetate, vitamin C palmitate, and the like; antioxidants including beta carotene, alpha hydroxy acids such as

glycolic acid, citric acid, lactic acid, malic acid, mandelic acid, ascorbic acid, alpha-hydroxybutyric acid, alpha-hydroxyisobutyric acid, alpha-hydroxyisocaproic acid, atrolactic acid, alpha-hydroxyisovaleric acid, ethyl pyruvate, galacturonic acid, glucoheptonic acid, glucoheptono 1,4-lactone, gluconic acid, gluconolactone, glucuronic acid, glucuronolactone, glycolic acid, isopropyl pyruvate, methyl pyruvate, mucic acid, pyruvic acid, saccharic acid, saccaric acid 1,4-lactone, tartaric acid, and tartronic acid; beta hydroxy acids such as beta-hydroxybutyric acid, beta-phenyl-lactic acid, beta-phenylpyruvic acid; botanical extracts such as green tea, soy, milk thistle, algae, aloe, angelica, bitter orange, coffee, goldthread, grapefruit, hoellen, honeysuckle, Job's tears, lithospermum, mulberry, peony, pueraria, nice, safflower, and mixtures thereof.

Preferred anti-aging agents include retinoids, anti-oxidants, alpha-hydroxy acids and beta-hydroxy acid with retinol and tretinoin being most preferred.

Suitable amounts of anti-aging agents include, based upon the total weight of the described cleaning system, from about 0.01 percent to about 20 percent, and preferably from about 0.04 percent to about 5 percent.

Examples of suitable anti-acne agents include, but are not limited to topical retinoids (tretinoin, isotretinoin, motretinide, adapalene, tazarotene, azelaic acid, retinol); salicylic acid; benzoyl peroxide; resorcinol; antibiotics such as tetracycline and isomers thereof, erythromycin, and the anti-inflammatory agents such as ibuprofen, naproxen, hetprofen; botanical extracts such as alnus, arnica, artemisia capillaris, asiasarum root, birrh, calendula, chamomile, cnidium, comfrey, fennel, galla rhois, hawthorn, houttuynia, hypericum, jujube, kiwi, licorice, magnolia, olive, peppermint, philodendron, salvia, sasa albo-marginata; imidazoles such as ketoconazole and elubiol, and those described in Gollnick, H et al. 196(l) Dermatology Sebaceous Glands, Acne and Related Disorders, 119-157 (1998), which is incorporated by reference herein, and mixtures thereof.

Preferred anti-acne agents include benzoyl peroxide, retinol, elubiol, antibiotics, and salicylic acid, with retinol and tretinoin being most preferred.

Suitable amount of anti-acne agents include, based upon the total weight of the described cleaning system, from about 0.01 percent to about 10 percent, and preferably from about 0.04 percent to about 5 percent.

Another preferred embodiment of the present invention is directed to a method for depigmenting the skin, comprising topically applying to skin at a desired area the above-described cleaning system and an effective amount of the depigmentation benefit agent. Suitable effective amounts of depigmentation agents include, based upon the total weight of the described cleaning system, from about 0.01 percent to about 10 percent, and preferably from about 0.04 percent to about 5 percent.

Examples of suitable depigmentation agents include, but are not limited to soy and derivatives thereof, retinoids such as retinol; Kojic acid and its derivatives such as, for example, kojic dipalmitate; hydroquinone and its derivatives such as arbutin; transexamic acid; vitamins such as niacin, vitamin C and its derivatives; azelaic acid; placertia; licorice; 5 extracts such as chamomile and green tea, and mixtures thereof, with retinol, Kojic acid, and hydroquinone, being preferred.

An alternative preferred embodiment of the present invention is directed to a method for treating the symptoms and/or the diseases of dandruff, seborrheic dermatitis and/or psoriasis, comprising topically applying the above-described cleaning system and the 10 relevant benefit agent to a location desired wherein the benefit agent is comprised of an effective amount of a dandruff treatment agent, a seborrheic dermatitis treatment agent, or a psoriasis treatment agent, respectively. As used herein, "dandruff treatment agent," "seborrheic dermatitis treatment agent," or a "psoriasis treatment agent," respectively, shall include agents capable of treating the symptoms and/or the diseases of dandruff, seborrheic 15 dermatitis, and psoriasis, respectively. By "effective amount," it is meant an amount effective for treating the disease and/or the symptoms associated therewith and preferably may range from, based upon the total weight of the cleaning system, from about 0.001 percent to about 10 percent, and preferably from about 0.01 percent to about 5 percent.

Examples of benefit agents suitable for treating the symptoms and/or the diseases 20 of dandruff, seborrheic dermatitis and/or psoriasis, respectively, nonexclusively include those set forth above with shale oil and derivatives thereof, elubiol, ketoconazole, coal tar, salicylic acid, zinc pyrithione, selenium sulfide, hydrocortisone, sulfur, menthol, pramoxine hydrochloride, and mixtures thereof being particularly preferred.

The compositions of the present invention may be directed applied to the skin or may 25 be applied onto other delivery implements such as wipes, sponges, brushes, and the like. The compositions may be used in products designed to be left on the skin, wiped from the skin, or rinsed off of the skin.

The invention illustratively disclosed herein suitably may be practiced in the absence of any component, ingredient, or step which is not specifically disclosed herein. Several 30 examples are set forth below to further illustrate the nature of the invention and the manner of carrying it out. However, the invention should not be considered as being limited to the details thereof.

Examples

Example 1: Preparation of Oil in Water Emulsion Incorporating a Polymeric Emulsifier**Preparation of Lipophilic Phase:**

11 g of PEG-30 dipolyhydroxystearate, available from Uniqema, Inc. under the tradename "Arlacel P-135," 50 g of isononyl isononanoate, available from Alzo, Inc. under the tradename, "Wickenol 151," 50 g of a mixture of hexyldecyl benzoate and butyloctyl benzoate, available from C.P. Hall Company under the tradename, "Hallstar AB," and 50 g of cyclomethicone available from Dow Corning under the tradename, "Dow 344 Fluid" were combined into a glass beaker containing a propeller stirrer and heated to a temperature of 60 °C until homogeneous.

Preparation of Hydrophilic Phase:

Into a primary glass beaker containing 795 g of deionized water, 5 g of PEG-8 caprylic/capric glycerides available from Trivent Inc. under the tradename, "Trivasol BW" was added thereto with stirring at 25°C until homogeneous. For aiding in dispersion of the thickener in the formulation, 4 g of carbomer available from B.F. Goodrich, Inc. under the tradename, "Carbopol Ultrez" were added to 30 g of dimethylisorbide available from Uniqema, Inc. under the tradename, "Arlasolve DMI" in a separate beaker with hand stirring. Into the dimethylisorbide mixture was then added 2 g. of methylparaben and 1 g of propylparaben with hand stirring until homogeneous to produce a pre-mixture. The pre-mixture was then added to the primary glass beaker with constant stirring until the resulting mixture was homogeneous.

Preparation of Final Composition:

The lipophilic phase was then added to the hydrophilic phase with constant stirring at 25°C until homogeneous. 2 g of triethanolamine available from Union Carbide under the tradename, "Trolamine 99%" was then added to the resulting mixture with stirring until homogeneous.

Example 2: Preparation of Polymeric Emulsifier-Free Oil in Water Emulsion**Preparation of Lipophilic Phase:**

20 g of isostearyl palmitate, available from Brooks Industries, under the tradename "Loronate OP," 20 g of isononyl isononanoate, available from Alzo, Inc. under the tradename, "Wickenol 151," 20 g of cetyl octanoate, available from Brooks Industries, under the tradename "Loronate CIO," 20 g of pentaerythritol tetraoctanoate available from Brooks Industries, under
 5 the tradename "Loronate PT," and 20 g of cyclomethicone available from Dow Corning under the tradename, "Dow 345 Fluid" were combined into a glass beaker at a temperature of 25 °C and stirred until homogeneous.

Preparation of Hydrophilic Phase:

Into a primary glass beaker containing 859.7 g of deionized water, 5 g of carbomer
 10 available from B.F. Goodrich, Inc. under the tradename, "Carbopol Ultrez" was added thereto with stirring at a temperature of 25 °C until homogenous. Into a separate beaker was added 7.5 g. of sucrose cocoate available from Croda, Inc. under the tradename, "Crodesta SL-40," 7.5 g. of PEG -6 Capric/caprylic glycerides available from Croda, Inc. under the tradename, "Glycerox 767," 10 g of hexylene glycol, 3 g. of methylparaben and 0.5 g of propylparaben with
 15 hand stirring until homogeneous to produce a pre-mixture. The pre-mixture was then added to the primary glass beaker with constant stirring until the resulting mixture was homogeneous.

Preparation of Final Composition:

After the lipophilic phase was added to the hydrophilic phase with constant stirring at 25°C until homogeneous, 6.8 g. of a 20% aqueous solution of sodium hydroxide was then
 20 added thereto with stirring at a temperature of 25 °C. The resulting mixture was then mixed for 15 minutes.

Example 3: Use of Emulsions to Remove Make-Up

Onto 3 respective ½ inch by 1 inch rectangle areas located on the forearms of
 25 caucasian women were applied each of the make-ups described in Table A below:

Table A: Types of Make-Up

<u>Type of Make-up</u>	<u>Supplier</u>	<u>Tradename / Color</u>
Foundation	Revlon	"Colorstay Make-Up" – in "Mocha"

Lipstick	Estee Lauder	"All Day Lipstick" – in "Regal Red"
Mascara	Revlon	"Colorstay Lashcolor Waterproof" - in "Navy"

After dipping a wooden applicator from Scientific Products into the foundation, the foundation was spread throughout the designated area and evenly distributed therein with a finger cot.

- 5 The lipstick was directly applied to the designated area and evenly distributed therein with a finger cot.

The mascara was spread with its wand in the designated rectangular area and evenly distributed therein with a finger cot.

- 10 After the make-up dried for 30 minutes at room temperature, 0.8 ml of the emulsion of Example 1 was dispensed via syringe across all three test sites. Using a wooden tongue depressor, the emulsion was rubbed across all three test sites in a rapid, circular motion for 30 seconds. Then, each of the three test sites were gently wiped with a Kim-wipe.

- This procedure was repeated but wherein the emulsion was replaced with the following respective cleansing products: a) "Galenic," which is available from Laboratoires Galenic, a
15 division of PIERRE FABRE, under the tradename, "Lait Demaquillant Hydratant (Moisturizing Cleansing Milk);" b) "Cetaphil," which is available from GALDERMA INTERNATIONAL under the tradename, "Lotion Nettoyante/Haute Tolerance (pour peaux sensibles);" c) "pH 5.5 3-in-1," which is available from JOHNSON & JOHNSON LIMITED under the tradename,
"JOHNSON's pH 5.5 – 3 in 1 Facial Cleanser;" d) "Vichy," which is available from VICHY
20 LABORATORIES under the tradename, "Demaquillant Integral (a l'eau thermale apaisante);" e) "Oil of Olay," which is commercially available from THE PROCTER & GAMBLE COMPANY under the tradename, "Oil of Olay Facial Cleansing Lotion;" and f) "F # 8626-015," which is similar to the formulation as set forth in Example 2, except as follows in Table B below:

Table B: Contrast of Example 2 formulation with formula 8626-015:

Example 2 Formulation	Modified Example 2 (F # 8626-015)	Component
0.3% (3 grams)	0.2% (2 grams)	methylparaben
0.05% (0.5 grams)	0.1% (1 gram)	Propylparaben

85.97% (859.7 grams)	86.02% (860.2 grams)	Water
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Photographs of the test sites were obtained after the make-up was applied and after the three sites were rubbed with the above six products as shown in FIGURE 1.

As evidenced in FIGURE 1, the emulsion of modified Example 2 was superior with respect to removal of mascara, foundation, and lipstick in comparison to the other cleansers. The emulsion of Example 1 (not shown in FIGURE 1) also was effective in removing all three types of make-up. This Example showed that the emulsions of the present invention are superior with respect to known commercial cleansers in removing a variety of make-up types from the skin.

Example 4: Preparation of Shower Gel Containing Sunscreen

A composition containing the following ingredients as set forth in Table C is prepared as follows:

Table C: Composition of Shower Gel Containing Sunscreen

<u>Tradename</u>	<u>Chemical Name</u>	<u>Weight (g)</u>	<u>Supplier</u>
WATER PHASE			
-----	Deionized water	696.5	-----
-----	Methylparaben	3	-----
Comperian 100K	Cocamide MEA	30	Henkel Corporation
Planteren 1200N	Lauryl glucoside	50	Henkel Corporation
Schercomid HT 60	PEG-50-Tallow Amide	50	Scher Chemical Corporation
	Hexylene glycol	10	
Standamox CAW	Cocamidopropylamine oxide	50	Henkel Corporation
OIL PHASE			
Hallstar AB	Hexyldecyl benzoate & butyloctyl benzoate	50	C.P. Hall Company
Arlacel P-135	PEG-30 Dipolyhydroxystearate	30	Uniqema, Inc.
Neo Heliopan	Menthyl anthranilate	30	Haarmann & Reimer
POST ADDITION			
Crothix Liquid	PEG 150 pentaerythrityl tetrastearate & PEG-6 caprylic/capric glycerides & water	10	Croda Inc.
Kathon CG	Methylchloroisothiazolinone & methylisothiazolinone	0.5	Rohm & Haas

After heating the deionized water to a temperature of about 75 °C to about 80 °C in a beaker, the hexylene glycol followed by the methylparaben is added thereto with mixing until solubilized. The cocamide MEA followed by the lauryl glucoside is then added thereto sequentially at constant temperature. The PEG-50 tallow amide is then melted and added to the resulting mixture at constant temperature. The cocamidopropylamine oxide is then added thereto at constant temperature to form a water phase premixture.

In another beaker, all of the oil phase components are combined with mixing at a temperature of about 75 °C to about 80 °C until uniform to form an oil phase premixture. The oil phase premixture is then added to the water phase premixture at constant temperature. The "Crothix Liquid" component is then added thereto with mixing until uniform, then the temperature is lowered to about 50 °C. The "Kathon CG" component is then added thereto, and the resulting mixture is cooled to room temperature.

The resulting cleansing formulation will possess excellent foaming properties and, after rinsing, will leave the skin with a light, moisturized feeling. Moreover, an effective amount of sunscreen agent will be deposited on the skin after rinsing the formulation from the skin with water.

Examples 5 and 6: Preparation of Foaming Anti-Acne Shower Gels

The composition of Example 5 containing the following ingredients as set forth in Table D is prepared as follows:

Table D: Compositions of Foaming Shower Gel

Tradename	Chemical Name	Wt % Ex. 5	Wt % Ex. 6	Supplier
WATER PHASE				
-----	Deionized water	49.440	49.654	-----
-----	Methylparaben	0.3	0.3	-----
Comperian 100K	Cocamide MEA	3.0	3.0	Henkel Corporation
Planteren 1200N	Lauryl glucoside	5.0	5.0	Henkel Corporation
Schercomid HT 50	PEG-50 Tallow Amide	5.0	5.0	Scher Chemical Co.
Standamox CAW	Cocamidopropylamine oxide	5.0	5.0	Henkel Corporation
Schercoquat DAS	Quaternium 61	2.0	2.0	Scher Chemical Co.
Jaguar C17	Guar hydroxypropyl trimonium chloride	0.5	0.5	Rhodia

	Glycerin	5.0	5.0	----
	Deionized water	10	10	----
	Citric acid (20% soln)	0.566	0.490	----
OIL PHASE				
Abil EM 90	Cetyl dimethicone copolyol	0.0	3.0	Goldschmidt Chemical Corporation
Arlacel P-135	PEG-30 Dipolyhydroxystearate	3.0	0.0	Unichema, Inc.
Wickenol 151	Isononyl isononanoate	1.0	1.0	Alzo
Volpo L3 Special	Laureth - 3	5.0	5.0	Croda, Inc.
POST ADDITION				
Arlasolve DMI	Dimethyl isosorbide	4.0	4.0	Croda Inc.
Salicylic acid	Salicylic acid	2.0	2.0	----
NaOH (10%)	Sodium hydroxide	0.144	none	----
Kathon CG	Methyl-chloroisothiazolinone & methylisothiazolinone	0.050	0.050	Rohm & Haas

Preparation of Hydrophilic Phase

The water phase premixture is prepared in accordance with Example 3, but with the following additional steps: The quaternium-61 is added with mixing to the mixture after the
5 addition of the cocamidopropyl amine oxide thereto at constant temperature until homogeneous.

In a separate glass beaker equipped with a propeller, the guar hydroxypropyl trimonium chloride and the glycerin are combined with stirring. The additional deionized water is then slowly added thereto with constant stirring. The resulting mixture is then mixed for
10 about 15 minutes, then acidified with the citric acid solution. The acidified solution is then added to the water phase premixture with stirring at a temperature of about 75 °C to about 80 °C until homogeneous.

Preparation of Oil Phase

PEG-30 dipolyhydroxystearate is combined with the laureth-3 in another beaker with
15 mixing at a temperature of about 75 °C to about 80 °C to form an oil phase premixture.

The oil phase premixture is then added to the water phase premixture with mixing at constant temperature.

Preparation of Final Composition:

The lipophilic phase is added to the hydrophilic phase with constant stirring at 75°C
20 until homogenous. After the temperature of the resulting mixture is lowered to 50°C with

continuous stirring, a premixed solution of 40 grams of Dimethyl Isosorbide available from Uniqema Inc. under the tradename "Arlasolve DMI" and 20 grams of salicylic acid are added thereto with stirring. After the temperature of the resulting mixture is lowered to 30°C, its pH is adjusted to approximately 3.1 by adding 1.4 grams of a 10% aqueous sodium hydroxide solution with continued stirring until the temperature of the resulting mixture reaches 25°C.

This process is repeated for the formulation of Example 6, but the PEG-30 dipolyhydroxystearate is replaced with cetyl dimethicone copolyol.

The above formulations will result in gels that possess good foaming properties as well as leave the skin with a pleasant, lightly moisturized "after feel" due to the effective deposition of the humectant. Similarly, the formulations will be useful as anti-acne formulations due to their ability to effectively deposit the salicylic acid component into and onto the skin.

Example 7: Preparation of Foaming Gel Containing Sunscreen

Preparation of Hydrophilic Phase:

Into a primary glass beaker equipped with a propeller containing 667 grams of deionized water, 30 grams of Hexylene Glycol available from Shell Chemical Company and 3 grams of methylparaben are added thereto stirring at 75°C until homogenous. While maintaining the agitation and temperature as constant, 30 grams of cocomonoethanolamide (100% active) available from Henkel Corporation under the tradename "Comperlan 100K", 50 grams of Lauryl Glucoside (50% active) available from Henkel Corporation under the tradename "Plantaren 1200N", 50 grams of PEG-50 Tallow Amide (100% active) available from Scher Chemical Corporation under the tradename "Schercomid HT 60", and 50 grams of Cocamidopropylamine Oxide (30% active) available from Henkel Corporation under the tradename "Standamox CAW" are added to this aqueous mixture and mixed until homogenous.

Preparation of Lipophilic Phase:

50 grams of Hexyldecyl Benzoate and Butyloctyl Benzoate available from C.P. Hall Co. under the tradename "HallStar AB", 30 grams of PEG-30 Dipolyhydroxystearate available from Unichema, Inc. under the tradename "Arlacel P135" and 30 grams of Menthyl Anthranilate available from Haarmann & Reimer Corp. under the tradename "Neo Heliopan MA" are combined into another glass beaker equipped with a propeller and stirred until homogenous with heat until the resulting lipophilic mixture was at a temperature of 75°C.

Preparation of Final Composition:

The lipophilic phase is then added to the hydrophilic phase with constant stirring at 75°C. 10 grams of a mixture of PEG-150 Pentaerythrityl Tetrastearate, PEG-6 Caprylic/Capric Glycerides, and Water available from Croda Inc. under the tradename "Crothix Liquid" is then added thereto. The resulting mixture is then stirred at 75°C until homogenous, then cooled to room temperature with constant stirring.

The resulting cleansing formulation possesses excellent foaming properties and, after rinsing, leaves the skin with a light, moisturized feeling. The formulation also effectively deposits the sunscreen agent into and onto the skin.

10

Example 8: Preparation of Foaming Acne Wash

Preparation of Hydrophilic Phase:

Into a primary glass beaker equipped with a propeller stirrer containing 414.9 grams of deionized water, 3 grams of methylparaben are added thereto at 75°C until homogenous. While maintaining constant agitation and temperature, 30 grams of cocomonoeethanolamide (100% active) available from Henkel Corporation under the tradename "Comperlan 100K", 50 grams of Lauryl Glucoside (50% active) available from Henkel Corporation under the tradename "Plantaren 1200N", 50 grams of PEG-50 Tallow Amide (100% active) available from Scher Chemical Corporation under the tradename "Schercomid HT 60", 50 grams of Cocamidopropylamine Oxide (30% active) available from Henkel Corporation under the tradename "Standamox CAW", and 20 grams of Quaternium-61 (90% active) available from Scher Chemical Corporation under the tradename "Schercoquat DAS" are added to this aqueous mixture and mixed until homogenous to form a primary hydrophilic phase.

5 grams of Guar Hydroxypropyl Trimonium Chloride available from Rhodia Inc. under the tradename "Jaguar C17" is combined with stirring with 50 grams of glycerin into a separate glass beaker equipped with a propeller stirrer. 100 grams of deionized water is slowly added with mixing to this mixture at constant temperature. After the resulting mixture is mixed for 15 minutes until homogenous, the mixture is acidified with 5.7 grams of a 20% aqueous solution of citric acid. After the acidified mixture is added to the primary hydrophilic phase, the resulting hydrophilic mixture is stirred at 75°C until homogenous.

Preparation of Lipophilic Phase:

50 grams of Isopropyl PPG-2-Isodeceth-7 Carboxylate available from Clariant Corporation under the tradename "Velsan D8P-3", 30 grams of PEG-30 Dipolyhydroxystearate available from Unichema, Inc. under the tradename "Arlacel P135" and 30 grams of Laureth-3 (100% active) available from Croda Inc. are combined with

stirring into a glass beaker equipped with a propeller stirrer until homogenous and heated to a temperature of 75°C.

Preparation of Final Composition:

- 5 The lipophilic phase is added to the hydrophilic phase with constant stirring at 75°C until homogenous. After the temperature of the resulting mixture is lowered to 50°C with continuous stirring, a premixed solution of 40 grams of Dimethyl Isosorbide available from Unichema Inc. under the tradename "Arlasolve DMI" and 20 grams of salicylic acid is added thereto with stirring. After the temperature of the resulting mixture is lowered to 30°C, its pH
- 10 is adjusted to approximately 3.1 by adding 1.4 grams of a 10% aqueous sodium hydroxide solution with continued stirring until the temperature of the resulting mixture reaches 25°C.

The resulting wash product will not only possess excellent foaming properties but will also be an effective anti-acne product due to its ability to deposit the anti-active agent into and onto the skin.

15

Example 9: Preparation of Water-in-Oil Emulsion

Preparation of Lipophilic Phase:

- 20 g of isostearyl palmitate, available from Brooks Industries, under the tradename
- 20 "Loronate OP," 20 g of isononyl isononanoate, available from Alzo, Inc. under the tradename, "Wickenol 151," 20 g of cetyl octanoate, available from Brooks Industries, under the tradename "Loronate CIO," 20 g of pentaerythritol tetraoctanoate available from Brooks Industries, under the tradename "Loronate PT," and 20 g of cyclomethicone available from Dow Corning under the tradename, "Dow 345 Fluid" were combined into a glass beaker at a temperature of 25 °C
- 25 and stirred until homogeneous.

Preparation of Hydrophilic Phase:

- Into a primary glass beaker containing 859.7 g of deionized water, 5 g of carbomer available from B.F. Goodrich, Inc. under the tradename, "Carbopol Ultrez" was added thereto with stirring at a temperature of 25 °C until homogenous. Into a separate beaker was added
- 30 7.5 g. of sucrose cocoate available from Croda, Inc. under the tradename, "Crodesta SL-40," 7.5 g. of PEG -6 Capric/caprylic glycerides available from Croda, Inc. under the tradename,

"Glycerox 767," 10 g of hexylene glycol, 3 g. of methylparaben and 0.5 g of propylparaben with hand stirring until homogeneous to produce a pre-mixture. The pre-mixture was then added to the primary glass beaker with constant stirring until the resulting mixture was homogeneous.

Preparation of Final Composition:

- 5 After the 6.8 g. of a 20% aqueous solution of sodium hydroxide was added to the hydrophilic phase with constant stirring at 25°C until homogeneous, the lipophilic phase was added thereto with stirring at a temperature of 25 °C. The resulting mixture was then mixed for 15 minutes.

Example 10: Preparation of Oil-in-Water Emulsion Containing Retinol

10 Preparation of Lipophilic Phase:

- 11 g of PEG-30 dipolyhydroxystearate, available from Uniqema, Inc. under the tradename "Arlacel P-135," 50 g of isononyl isononanoate, available from Alzo, Inc. under the tradename, "Wickenol 151," and 50 g of a mixture of hexyldecyl benzoate and butyloctyl benzoate, available from C.P. Hall Company under the tradename, "Hallstar AB" were
15 combined with continuous mixing in a vessel and heated to a temperature of 45°C until homogeneous. After the mixture was cooled to a temperature of 25 °C, 50 g of cyclomethicone available from Dow Corning under the tradename, "Dow 344 Fluid" and 6.9 g of a mixture of vitamin A alcohol and polysorbate 20 in a 1:1 weight ratio were added thereto with continuous mixing under an Argon blanket and under yellow light into a glass beaker containing a propeller
20 stirrer until homogeneous. All subsequent procedures with this lipophilic phase were conducted under these conditions of argon blanket and yellow light until the formulation is placed into an oxygen and light impermeable container.

Preparation of Hydrophilic Phase:

- Into a primary glass beaker containing 795 g of deionized water, nitrogen was bubbled
25 therein until the subsequent addition of the lipophilic phase thereto so as to minimize exposure to oxygen. 5 g of PEG-8 caprylic/capric glycerides available from Trivent Inc. under the tradename, "Trivasol BW" was then added thereto with stirring at 25°C until homogeneous. For aiding in dispersion of the thickener in the formulation, 4 g of carbomer available from B.F.

- Goodrich, Inc. under the tradename, "Carbopol Ultrez" were added to 30 g of dimethylisosorbide available from Uniqema, Inc. under the tradename, "Arlasolve DMI" in a separate beaker with hand stirring. Into the dimethylisosorbide mixture was then added 2 g. of methylparaben and 1 g of propylparaben with hand stirring until homogeneous to produce a
- 5 pre-mixture. The pre-mixture was then added to the primary glass beaker with constant stirring until the resulting mixture was homogeneous.

Preparation of Final Composition:

- The lipophilic phase was then added to the hydrophilic phase with constant stirring at 25°C until homogeneous. 2 g of triethanolamine available from Union Carbide under the
- 10 tradename, "Trolamine 99%" was then added to the resulting mixture with stirring until homogeneous. The final emulsion contains the components as set forth in Table E:

Table E: Emulsion Components

Chemical Name	Trade Name	%(wt/wt)
PEG-30 dipolyhydroxystearate	Arlacel P-135	1.1
Isononyl isononanoate	Wickenol	5.0
Hexyldecyl benzoate and butyloctyl benzoate	Hallstar AB	5.0
Cyclomethicone	Dow 344 Fluid	5.0
Vitamin A alcohol and Tween 20	Retinol 50C	0.69
Water	Water	78.81
Carbomer	Carbopol Ultrez	0.40
PEG-8 caprylic/capric glycerides	Trivasol BW	0.50
Methylparaben	Methylparaben	0.20
Propylparaben	Propylparaben	0.10
Dimethyl isosorbide	Arlasolve DMI	3.0
triethanolamine	Trolamine 99%	0.2

Example 11: Preparation of Water-in-Oil Emulsion Containing Retinol

Preparation of Lipophilic Phase:

11 g of PEG-30 dipolyhydroxystearate, available from Uniqema, Inc. under the tradename "Arlacel P-135," 30 g of isononyl isononanoate, available from Alzo, Inc. under the tradename, "Wickenol 151," and 30 g of a mixture of hexyldecyl benzoate and butyloctyl benzoate, available from C.P. Hall Company under the tradename, "Hallstar AB" were combined in a vessel with mixing and heated to a temperature of 45 °C until homogeneous. After the resulting mixture was cooled to a temperature of 25°C, 30 g of cyclomethicone available from Dow Corning under the tradename, "Dow 344 Fluid" and 6.9 g of a mixture of vitamin A alcohol and polysorbate 20 in a 1:1 weight ratio were added thereto with continuous mixing under an Argon blanket and under yellow light into a glass beaker containing a propeller stirrer until homogeneous. All subsequent procedures with this lipophilic phase was conducted under these conditions of argon blanket and yellow light until the formulation is placed into an oxygen and light impermeable container.

Preparation of Hydrophilic Phase:

Into a primary glass beaker containing 863.2 g of deionized water, nitrogen was bubbled therein in order to eliminate dissolved oxygen contained therein. The nitrogen continued to be bubbled therein until the subsequent addition of the lipophilic phase thereto. 5 g of PEG-8 caprylic/capric glycerides available from Trivent Inc. under the tradename, "Trivasol BW" was then added thereto with stirring at 25°C until homogeneous. For aiding in dispersion of the thickener in the formulation, 4 g of carbomer available from B.F. Goodrich, Inc. under the tradename, "Carbopol Ultrez" were added to 10 g of triisopropyl citrate available from Phoenix Chemical Company under the tradename, "PELEMOL TIPC" in a separate beaker with hand stirring. Into the triisopropyl citrate mixture was then added 2 g. of methylparaben and 1 g of propylparaben with hand stirring until homogeneous to produce a pre-mixture. The pre-mixture was then added to the primary glass beaker with constant stirring until the resulting mixture was homogeneous.

Preparation of Final Composition:

2 g of triethanolamine available from Union Carbide under the tradename, "Trolamine 99%" was then added to the hydrophilic phase with constant stirring at 25°C until homogeneous. The resulting mixture was then added to the lipophilic phase at constant temperature with stirring until homogeneous. The final emulsion contains the components as set forth in Table F:

Table F: Emulsion Components

Chemical Name	Trade Name	%(wt/wt)
PEG-30 dipolyhydroxystearate	Arlacel P-135	1.1
Isononyl isononanoate	Wickenol	3.0
Hexyldecyl benzoate and butyloctyl benzoate	Hallstar AB	3.0
Cyclomethicone	Dow 344 Fluid	3.0
Vitamin A alcohol and Tween 20	Retinol 50C	0.69
Water	Water	86.320
Carbomer	Carbopol Ultrez	0.40
PEG-8 caprylic/capric glycerides	Trivasol BW	1.0
Methylparaben	Methylparaben	0.20
Propylparaben	Propylparaben	0.10
Triisopropyl citrate	Pelemol TIPC	1.0
NaOH	NaOH	0.190

Example 12 – Luminosity of the Formulation of Example 10

Digital images of the right side and the left side of a Caucasian woman's face was taken using a digital camera available from Fujix (Model No.: DCS 505) equipped with a 60 mm macro lens under strobe light conditions at F8 and 1/125 seconds. The camera lens was filtered with a CG-395 filter, and the strobe light source was filtered with a combination of a UG-11 filter and a KG-5 filter. These images are illustrated in FIG. 3(a) and FIG. 3(b), respectively.

After approximately 0.09 grams of the 0.3% retinol formulation prepared in Example 10 was applied to about a 20 cm² site on the suborbital (cheek) area of the right side and the

left side of the woman's face, digital images were taken thereof under the above conditions as illustrated in FIG. 3(c) and FIG. 3(d), respectively. Using PHOTOSHOP software available from Adobe Inc., the digital image of each site was analyzed for average pixel intensity or luminosity. Luminosity, as used herein, is an indication of brightness of a given area as measured on a scale of 1 to 255, wherein the latter is the most luminescent. Using the 0.3% retinol concentration value, the pixel intensity change, as determined by the difference in pixel intensity between the base surface and the treated surface, for both the treated right side and left side of the face was plotted as a function thereof as illustrated in FIG. 5 (a).

The formulation was then rinsed from the right side of the face, and a digital image was taken of the site under the above conditions as illustrated in FIG. 3(e). The formulation was then wiped twice using a Kimwipe tissue available from Kimberly Clark from the left side of the face, and a digital image was taken of the site under the above conditions as illustrated in FIG. 3(f). The pixel intensity change for the rinsed right side and the wiped left side was plotted on the graph of FIG. 5(a), then the respective deposited retinol concentrations were interpolated therefrom to be 0.145% and 0.1%, respectively.

This Example showed that the formulation of the present invention is not only a cleanser, but it also effectively deposits active agents, such as retinol, onto the skin. A significant amount of the agents remained on the skin after the formulation was removed therefrom. Moreover, this Example highlighted that when the cleanser composition of the present invention contains a 0.3% retinol active agent, it deposited the same amount of retinol on the skin as a leave-on product containing 0.145% retinol (when the compositions was removed via rinsing with water) and a leave-on product containing a 0.1% retinol (when the composition was removed via wiping).

Example 13 – Luminosity of the Formulation of Example 11

The procedure set forth in Example 12 was repeated using the formulation of Example 11 instead of that of Example 10. The pre-treatment images are illustrated in FIG. 4(a) and FIG. 4(b), respectively.

The formulation-containing images are illustrated in FIG. 4(c)(right side) and FIG. 4(d)(left side). Using the 0.3% retinol concentration value, the pixel intensity change for the treated right side and left side of the face was plotted as a function thereof as illustrated in FIG. 5 (d).

The digital image of the washed site is illustrated in FIG. 4(e), and the image of the wiped side is illustrated in FIG. 4(f). The pixel intensity change for the rinsed right side and the wiped left side was plotted on the graph of FIG. 5(b), then the respective deposited retinol concentrations were interpolated therefrom to be 0.135% and 0.072%, respectively.

This Example showed that the formulation of the present invention was not only a cleanser, but it also effectively deposited active agents, such as retinol, onto the skin. These agents remained present on the skin after the formulation was removed therefrom.

Moreover, this Example highlighted that when the cleanser composition of the present

- 5 invention contained a 0.3% retinol active agent, the cleanser deposited the same amount of retinol on the skin as a leave-on product containing 0.135% retinol (when the compositions was removed via rinsing with water) and a leave-on product containing a 0.072% retinol (when the composition was removed via wiping).

10 **Example 14: Preparation of Oil-in-Water Emulsion Containing DMAE**

Preparation of Lipophilic Phase:

- 10 g of steareth-2 available from Uniqema under the tradename "Brij 72", 8.5 g of isoceteth-20 also available from Uniqema under the tradename "Arlasolve 200", 10 g. of isononyl isononanoate available from Alzo, Inc., under the tradename "Wickenol 151", 10 g. of
15 Isostearyl palmitate available from Brooks Industries under the tradename "Loronate OP", 10 g of cetyl octanoate, available from Brooks Industries, under the tradename "Loronate CIO," 10 g of pentaerythritol tetraoctanoate also available from Brooks Industries, under the tradename "Loronate PT," and 10 g of cyclomethicone available from Dow Corning under the tradename, "Dow 345 Fluid" were combined into a glass beaker at a temperature of 50 °C and stirred until
20 homogeneous.

Preparation of Hydrophilic Phase:

- 602.5 g of deionized water were weighed into a primary glass beaker and heated to 78-82°C. With constant agitation, 4 g of PVM/MA Decadiene Crosspolymer available from ISP under the tradename, "Stabileze QM" was added thereto and held at 78-82°C until
25 homogenous. This mixture was then cooled to 40-50°C, during which time, 1 g. of disodium EDTA, 10 g. of hexylene glycol, 7.5 g. of PEG-6 caprylic/capric glycerides available from Croda, Inc. under the tradename, "Glycerox 767", and 10 g. of PEG-150 pentaerythrityl tetrastearate also available from Croda under the tradename, "Crothix Liquid," were added to the primary beaker with constant stirring.

- 30 Preparation of Final Composition:

When both the lipophilic phase and the hydrophilic phase were at a temperature of 40°C -50°C, the lipophilic phase was added to the hydrophilic phase with constant stirring. In a separate beaker, 30 g. of 2-(dimethylamino) ethanol, available from BASF under the tradename DMAE, and 50 g. of L-tyrosine available from Ajinomoto under the tradename "L-Tyrosine" were added to 150 g. of water, and mixed until homogenous. This premix was then added to the primary beaker with constant stirring. 10 g of Nylon-12 available from Kobo Products, Inc., under the tradename "SP-10", 5 g of talc available from Luzenac America under the tradename, "Windsor Talc 66", 10 g of silicone quaternium-13 available from Biosil Industries under the tradename, "Biosil Basics SPQ," and 10 grams of a phenoxyethanol, methylparaben, butylparaben, ethylparaben and propylparaben solution available from Nipa under the tradename "Phenonip" were added separately to the primary beaker with constant stirring. The entire mixture was adjusted to a pH of 7.0-7.5 with a 70% aqueous solution of glycolic acid, and homogenized for 2 minutes at medium power with a Gifford-Wood homogenizer.

After about 1 ml to about 10 ml of the resulting formulation is applied to the facial skin of consumers, the consumers perceive that their facial skin appears and feels firmer and "lifted."

Example 15: *Preparation of Oil in Water Emulsion Containing a Polymeric Emulsifier and Colloidal Oat Flour*

Preparation of Hydrophilic Phase:

Into a primary glass beaker containing 850.70 g of deionized water, 10g of Colloidal Oat Flour available from Quaker was added thereto with stirring at 25°C until a homogeneous, smooth slurry was achieved. 2.5 g. of Acrylates/C10-30 Alkyl Acrylate Crosspolymer available from B.F. Goodrich, Inc. under the tradename, "Pemulen TR-1" and 2.5 g. of Carbomer, also available from B.F. Goodrich, Inc. under the tradename "Carbopol Ultrez" were then added to the primary beaker and mixed with slower agitation until homogenous. Into a separate beaker was added 7.5 g. of sucrose cocoate available from Croda, Inc. under the tradename, "Crodesta SL-40," 7.5 g. of PEG -6 Capric/caprylic glycerides available from Croda, Inc. under

the tradename, "Glycerox 767," 10 g of hexylene glycol, 3 g. of methylparaben and 0.5 g of propylparaben with hand stirring until homogeneous to produce a pre-mixture. The pre-mixture was then added to the primary glass beaker with constant stirring until the resulting mixture was homogeneous.

5 Preparation of Final Composition:

20 g of isostearyl palmitate, available from Brooks Industries, under the tradename "Loronate OP," 20 g of isononyl isononanoate, available from Alzo, Inc. under the tradename, "Wickenol 151," 20 g of cetyl octanoate, available from Brooks Industries, under the tradename "Loronate CIO," 20 g of pentaerythritol tetraoctanoate available from Brooks Industries, under
10 the tradename "Loronate PT," and 20 g of cyclomethicone available from Dow Corning under the tradename, "Dow 345 Fluid" were each added separately to the primary beaker with constant stirring at 25°C until homogeneous. 2.5 g of Tetrasodium EDTA and , 6.8 g. of a 20% aqueous solution of sodium hydroxide was then added thereto with stirring at a temperature of 25 °C. The resulting mixture was then mixed for 15 minutes.

15

**Example 16: Preparation of Oil in Water Emulsion
Containing a Polymeric Emulsifier and Colloidal Oat Flour**

20 Preparation of Preservative Pre-Blend

4 g of methylparaben, 1 g of propylparaben, 7.5 g of PEG-6 capric/caprylic glycerides available from Croda, Inc. under the tradename, "Glycerox 767," 7.5 g of sucrose cocoate also available from Croda, Inc. under the tradename, "Crodesta SL-40," and 10 g of hexylene glycol were combined with mixing under ambient conditions until homogeneous.

25

Preparation of Emulsion:

Into a primary glass beaker containing 852.5 g of Purified Water (USP), 10g of Colloidal Oat Flour available from Quaker were added thereto with stirring at about 200 rpm and a temperature of about 20 °C to about 30 °C until a homogeneous, smooth slurry was
30 achieved. 2.5 g. of Acrylates/C10-30 Alkyl Acrylate Crosspolymer available from B.F. Goodrich, Inc. under the tradename, "Pemulen TR-1" and 2.5 g. of Carbomer, also available

from B.F. Goodrich, Inc. under the tradename "Carbopol Ultrez" were then added thereto and mixed with slower agitation until homogenous. After adding the Preservative Pre-blend with increased mixing at about 200 rpm thereto, the following components were sequentially added thereto with constant stirring at about 20°C to about 30°C until homogeneous, with intervals of

5 5 minutes between the addition of each respective component: 20 g of isononyl isononanoate, available from Alzo, Inc. under the tradename, "Wickenol 151," 20 g of cyclomethicone available from Dow Corning under the tradename, "Dow 345 Fluid" , 20 g of isostearyl palmitate, available from Brooks Industries, under the tradename "Lorionate OP," 20 g of cetyl octanoate, available from Brooks Industries, under the tradename "Lorionate CIO," 20 g of

10 pentaerythritol tetraoctanoate available from Brooks Industries, under the tradename "Lorionate PT." 2.5 g of Tetrasodium EDTA and enough of a 20% aqueous solution of sodium hydroxide was then added thereto with stirring at a temperature of about 20°C to about 30°C to produce a final mixture having a pH of 5.9 to 6.5. The resulting mixture was then mixed until homogeneous.

15

Example 17– Consumer Testing of Formulation of Example 16 Formula

One hundred and twenty-five mothers of babies aged 24 months and younger used both the formulation prepared in accordance with Example 16 as well as a cleanser available from Galderma Laboratories, Inc. under the tradename, "CetaPhil Gentle Skin Cleanser."

20 The mothers used each product for a minimum of at least 3 to 7 times for a one week period. When using the product in a rinse-off fashion, the mothers first poured the product onto a moistened hand or wet cloth then applied the product to the desired location on the babies' skin. After rubbing the product gently on the skin, the product was rinsed therefrom with water. When using the product in a wipe-off fashion, the mothers applied a liberal amount to the

25 desired location on the babies' skin and rubbed gently. The excess product was then removed therefrom with a soft cloth or tissue.

The results of the study are shown in Table G below:

Table G Comparative Study of Example 16 Formulation to Cetaphil

Characteristic that either Completely describes or very well describes the product at issue	Formulation of Example 16	Cetaphil
1) Good for Sensitive Skin	87*	82
2) Nonirritating to Skin	92	91
3) Won't Dry or Irritate the Most Sensitive Skin	89	78
4) Cleanses Extra Gently	87	86
5) Good for Everyday or Regular Use	94	84
6) Cleans Without Drying Skin	92	85
7) Relieves Baby's Dry Skin	74	65
8) Can Be Use With or Without Water	93	83
9) Effectiveness as a Cleanser	82	76
10) Good for Use All Over the Body	90	85
11) Makes Baby's Skin Feel Soft and Smooth	89	74
12) Helps Baby's Skin Look and Feel Healthier	70	66
13) Helps Baby's Skin Retain its Natural Moisture	82	72
14) Relieves Baby's Itchy Skin	51	46
15) Leaves Baby's Skin Feeling Clean	86	75
16) Won't Sting or Irritate Eyes	54	54

*These numbers indicate the percentage of the mothers that indicated that the identified product possessed the given characteristic.

- This Example showed that the cleanser of Example 16 significantly outperformed the commercial product with respect to the majority of characteristics set forth in Table G.

Example 18 – Preparation of Oil-In-Water Emulsion with Non-Ionic Emulsifier

Preparation of Lipophilic Phase:

- 20 g of isostearyl palmitate, available from Brooks Industries, under the tradename "Loronate OP," 20 g of isononyl isononanoate, available from Alzo, Inc. under the tradename, "Wickenol 151," 20 g of cetyl octanoate, available from Brooks Industries, under the tradename

"Loronate CIO," 20 g of pentaerythritol tetraoctanoate available from Brooks Industries, under the tradename "Loronate PT," were combined into a glass beaker at a temperature of 25 °C and stirred until homogeneous.

Preparation of Hydrophilic Phase:

- 5 Into a primary glass beaker containing 859.7 g of deionized water, 2 g of carbomer available from B.F. Goodrich, Inc. under the tradename, "Carbopol ETD 2020" , and 1 g of C10-C30 alkyl acrylate/crosspolymer commercially available from B.F. Goodrich under the tradename, "Pemulen TR1" were added thereto with stirring at a temperature of 25 °C until dispersed. While heating the mixture to 75 °C, 1.2 g of tromethamine, 1g of EDTA, 7.5 g. of
- 10 PEG -6 Capric/caprylic glycerides available from Croda, Inc. under the tradename, "Glycerox 767," 10 g of hexylene glycol, 4 g. of methylparaben and 1 g of propylparaben were added with constant stirring until the resulting mixture was homogeneous. After the mixture reached a temperature of 75 °C, 10 g. of a mixture of sorbitan stearate and sucrose cocoate available from Uniqema under the tradename, "Arlatone 2121," were added thereto with stirring for 30
- 15 minutes at constant temperature.

Preparation of Final Composition:

- After the lipophilic phase was heated to a temperature of 75 °C, it was then added to the hydrophilic phase with constant stirring at 75°C until homogeneous. After the mixture was then cooled to 35 °C, 20 g of cyclomethicone available from Dow Corning under the
- 20 tradename, "Dow 345 Fluid" was added thereto. After the mixture was cooled to 25 °C, 0.4 g. of tromethamine was then added thereto with stirring at constant temperature such that the resulting mixture had a pH of 5.5.

Example 19: Comparison of Make-Up Removability

- 25 A waterproof mascara available from Gemey under the tradename "Waterproof Gemey Noir," is copiously coated on a 16 cm² area on a forearm. This area is then wiped four times with a cotton ball soaked with 2 ml of the formula of Example 2. Visual assessment of

the area is noted. This procedure is repeated on the same area and a second visual assessment is noted. This procedure is repeated on the same area and a third visual assessment is noted.

5 This procedure is repeated on an alternative areas of the forearm, but wherein the cleanser of Example 2 is replaced with the cleanser of Example 18 and the cleanser available from Johnson's under the tradename, "Johnson's pH 5.5 3-in-1 Cleansing Lotion," respectively.

Although all three cleansers effectively remove the mascara, this Example shows that the cleansers of Example 2 and Example 18 remove mascara more quickly than the commercial cleanser. In addition, all three cleansers are aesthetically acceptable according to
10 consumer standards.

Example 20: Comparison of Make-Up Removability of Impregnated Wipes

The formulation of Example 18 is prepared, but wherein: 1) the carbomer and the C10-C30 alkyl acrylate/crosspolymer were replaced with 1 g of xanthan gum; 2) and 30 g of
15 C14 – 22 alkyl alcohol and C12 –C20 alkyl glucoside emulsifier available from Seppic under the tradename "Montanov L" was also added to the lipophilic phase; and 3) and 4 g of C13-C14 isoparaffin/isostearyl isostearate/Na polyacrylate/polyacrylamide/polysorbate 60 available from Seppic under the tradename," Sepigel 502," was added to the finished mixture as the final product was cooled to 25 C.

20 The emulsion was pumped on to a stack of 25 folded, uncoated spunlaced wipes comprised of a blend of about 35% polyester and 65% rayon in an amount equivalent to 325% of the total noncoated wipe weight.

The resulting wipes were then compared with Johnson pH5.5 3-in-1 Cleansing wipes and Pond's Cleansing Towelettes available from Unilever using the test method set forth in
25 Example 19 wherein the cotton ball was replaced by the respective wipe.

This Example showed that the wipes prepared in accordance with this Example are more effective in removing waterproof mascara than the two commercial products as determined after the first, second, and third assessment. More specifically, the wipes of this

Example were significantly superior to the Pond's wipes with respect to mascara removal as determined after the third visual assessment.

Example 21: Preparation of Cleansing Composition

5

20 g of cyclomethicone available from the Dow Corning Corporation under the tradename, "DOW CORNING 345," 15 g of hexylene glycol, and 65 g of a mixture of hexyl decyl benzoate and butyl octyl benzoate available from the C.P. Hall Company under the tradename, "Hallstar AB" are sequentially added to a vessel with mixing at about 100 rpm
10 under ambient conditions until the final mixture is homogeneous.

The resulting cleansing composition is effective in removing debris such as makeup from the skin.

We claim:

1. A cleansing composition comprising
 - a. a liquid silicone;
 - 5 b. a water dispersible component and
 - c. an ester.
2. The composition of claim 1 wherein the composition is comprised of, based upon the total weight of the composition,
 - a. from about 10 percent to about 35 percent of the liquid silicone;
 - 10 b. from about 10 percent to about 35 percent of the water dispersible component;
 - and
 - c. from about 55 percent to about 65 percent of the ester.
3. The cleansing composition of claim 1 wherein the silicone is selected from the group consisting of hexamethylsiloxane, dimethicone, dimethiconol, cyclomethicone, and
15 mixtures thereof.
4. The cleansing composition of claim 3 wherein the cyclomethicone is selected from the group consisting of cyclo tetradimethyl siloxane; cyclopentadimethyl siloxane, cyclohexadimethyl siloxane, cycloheptadimethyl siloxane, and mixtures thereof.
5. The cleansing composition of claim 1 wherein the water dispersible component is
20 selected from the group consisting of polyethylene glycol 400, hexylene glycol, propylene glycol, polypropylene glycol-10 methylglucose ether, ethoxydiglycol, polyethylene glycol-6 caprylic/capric glycerides, ethylene glycol monobutyl ether, triisopropyl citrate, polyethylene glycol-8 caprylic/capric glycerides, 3-methoxy-3-methyl-1-butanol, dimethyl isosorbide, polyethylene-6 caprylic/capric triglyceride, and mixtures thereof.
6. The cleansing composition of claim 5 wherein the water dispersible component is
25 selected from the group consisting of hexylene glycol, dimethyl isosorbide, polyethylene glycol-6 caprylic/capric glyceride, and mixtures thereof.
7. The cleansing composition of claim 1 wherein the water dispersible component is comprised of, based upon the total weight percent of the cleaning composition,
30
 - a) from about 5 percent to about 15 percent of hexylene glycol;
 - b) from about 5 percent to about 10 percent of polyethylene-6 caprylic/capric triglyceride.

8. The cleansing composition of claim 1 wherein the ester is selected from liquid esters that either possess a structural means for ensuring the liquidity of the ester or are heterogeneous in nature.

5 9. The cleansing composition of claim 1 wherein the ester is selected from the group consisting of

a) a branched C₅ to C₂₂ alkyl alcohol ester of an aromatic acid;

b) a straight-chained or branched C₅ to C₂₂ alkyl acid esters of optionally ethoxylated/propoxylated polyols having from about 3 carbon atoms to about 7 carbon atoms;

10 c) branched C₅ to C₂₂ alkyl alcohol esters of branched polyacids;

d) branched or straight-chained C₅ to C₂₂ alkyl acid esters of branched and/or unsaturated C₅ to C₂₂ alkyl alcohols;

e) branched or unsaturated C₅ to C₂₂ alkyl alcohol esters of an acid selected from the group consisting of adipic acid, succinic acid, sebacic acid, maleic acid, and mixtures thereof

15 f) polyether interrupted fatty acid esters;

g) benzoic acid ester of heterogeneous alcohols having from about 8 carbon atoms to about 22 carbon atoms; and

h) mixtures thereof,

20 10. The cleansing composition of claim 9 wherein the ester is selected from the group consisting of straight-chained or branched C₅ to C₂₂ alkyl acid esters of optionally ethoxylated/propoxylated polyols; benzoic acid esters of heterogeneous alcohols; and mixtures thereof.

25 11. The cleansing composition of claim 9 wherein the ester is selected from the group consisting of butyloctyl salicylate; hexyldecyl benzoate; and butyloctyl benzoate; alkyl benzoates having from about 12 carbon atoms to about 15 carbon atoms; and mixtures thereof.

12. The cleansing composition of claim 11 wherein the ester is selected from the group consisting of hexyldecyl benzoate, butyloctyl benzoate, and mixtures thereof.

30 13. The cleansing composition of claim 9 wherein the ester is selected from the group consisting of pentaerythritol tetraoctanoate; trimethylolpropane trioctanoate; trioctanoin; pentaerythrityl tetrapelargonate; sorbitan trioleate; caprylic/capric triglyceride; neopentyl alcohol tetraoctanoate, and mixtures thereof.

14. The cleansing composition of claim 13 wherein the ester is selected from the group consisting of caprylic/capric triglyceride; pentaerythritol tetraoctanoate; trimethylolpropane trioctanoate; pentaerythrityl tetrapelargonate; and mixtures thereof.

5 15. The cleansing composition of claim 9 wherein the ester is selected from the group consisting of branched alkyl alcohol esters of branched polyacids, wherein the alkyl alcohol is optionally substituted and contains from about 3 carbon atoms to about 22 carbon atoms.

16. The cleansing composition of claim 15 wherein the ester is trioctyldecyl citrate and mixtures thereof.

10 17. The cleansing composition of claim 9 wherein the ester is selected from the group consisting of tridecyl neopentanoate, isostearyl palmitate, cetyl ricinoleate, cetyl octanoate, isononyl isononanoate, butyl stearate, octyldodecyl soyate, tridecyl erucate, octyldodecyl erucate/eicosil erucate, and mixtures thereof.

15 18. The cleansing composition of claim 17 wherein the ester is selected from the group consisting of cetyl octanoate, isostearyl palmitate, isononyl isononanoate, and mixtures thereof.

19. The cleansing composition of claim 9 wherein the ester is selected from the group consisting of diisopropyl adipate, dioctyl sebacate, dioctyl succinate, dioctyl maleate, diisostearyl adipate, diethyl sebacate, and mixtures thereof.

20 20. The cleansing composition of claim 19 wherein the ester is selected from the group consisting of diethyl sebacate, dioctyl sebacate, diisostearyl adipate, and mixtures thereof.

25 21. The cleansing composition of claim 9 wherein the ester is selected from the group consisting of laureth-2 benzoate; C_8 to C_{22} fatty alkyl (optionally polypropylenoxy) polyethyleneoxy carboxylate esters derived from an alcohol having from about 1 carbon atom to about 22 carbon atoms; and mixtures thereof.

22. The cleansing composition of claim 21 wherein the ester is isopropyl propylene glycol-2-isodeceth-7 carboxylate.

30 23. The cleansing composition of claim 9 wherein the ester is selected from at least two of the following esters:

a) branched C_5 to C_{22} alkyl alcohol esters of an aromatic acid;

b) branched or straight-chained C_5 to C_{22} alkyl acid esters of branched or unsaturated C_5 to C_{22} alkyl alcohols; and

c) straight-chained or branched C₅ to C₂₂ alkyl acid esters of optionally ethoxylated/propoxylated polyols.

23. The cleansing composition of claim 9 wherein the ester is a mixture of, based upon the total weight percent of the esters,:

5 a) from about 30 percent to about 80 percent of branched or straight-chained C₅ to C₂₂ alkyl acid esters of branched or unsaturated C₅ to C₂₂ alkyl alcohols;

 b) from about 10 percent to about 50 percent of branched C₅ to C₂₂ alkyl alcohol esters of an aromatic acid; and

10 c) from about 10 percent to about 50 percent of straight-chained or branched C₅ to C₂₂ alkyl acid esters of optionally ethoxylated/propoxylated polyols.

24. The cleansing composition of claim 9 wherein the ester is a mixture comprised of, based upon the total weight percent of the ester,:

 a) from about 15 percent to about 50 percent isononyl isononanoate;

 b) from about 15 percent to about 50 percent isostearyl palmitate;

15 c) from about 15 percent to about 50 percent cetyl octanoate; and

 d) from about 15 percent to about 50 percent pentaerthritol tetraoctanoate.

25. A cleaning system comprising:

 a. the cleansing system of claim 1;

 b. water; and

20 c. a polymeric emulsifier and/or a thickener.

26. The cleansing system of claim 25 comprising, based upon the total weight of the cleansing system:

 a. at least 5 percent of the cleansing system of claim 1;

 b. from about 70 percent to about 98 percent of water; and

25 c. from about 0.5 to about 1.5 percent of a polymeric emulsifier and/or thickener.

27. The cleansing system of claim 25 wherein the polymeric emulsifier is polyethylene glycol-30 dipolyhydroxystearate; dimethicone copolyol; substituted acrylates; and mixtures thereof.

30 28. The cleansing system of claim 25 wherein the thickener is selected from the group consisting of carbomers, acrylate copolymers, hydroxyethylcellulose modified with

cetyl ether groups, polyvinylmethyl ether/maleic anhydride (PVM/MA) decadiene crosspolymer, and mixtures thereof.

29. The cleansing system of claim 25 wherein the thickener is selected from the group consisting of acrylates/aminoacrylates copolymer, acrylates/steareth-20 methacrylate copolymer; acrylates/ceteth-20 itaconate copolymer, acrylates/steareth-20 itaconate copolymer, carbomers, modified hydroxycellulose, polyvinylacetate/maleic anhydride (PVA/MA) decadiene crosspolymer, and mixtures thereof.

30. The cleansing system of claim 25 further comprising, based upon the total weight of the cleansing system, from about 1 percent to about 3 percent of a cleansing enhancer.

31. The cleansing system of claim 30 wherein the cleansing enhancer is a nonfoaming surfactant and/or a non-ionic emulsifier.

32. The cleansing system of claim 31 wherein the nonfoaming surfactant is selected from the group consisting of sucrose cocoate, sucrose stearate and mixtures thereof.

33. The cleansing system of claim 31 wherein the non-ionic emulsifier is selected from the group consisting of isoceteth 20, oleth-2, mixture of PEG-40 hydrogenated castor oil and trideceth-9, Poloxamer 184, laureth-4, sorbitan trioleate, polyoxyethylene-(2) oleyl ether, sorbitan stearate, cetearyl glucoside, glyceryl oleate, and mixtures thereof.

34. The system of claim 25 further comprising a benefit agent.

35. The system of claim 34 wherein the benefit agent is selected from the group consisting of vasoconstrictors, collagen enhancers, anti-edema agents, depigmentation agents; reflectants; detangling/wet combing agents; film forming polymers; humectants; amino acid agents; antimicrobial agents; allergy inhibitors; anti-acne agents; anti-aging agents; anti-wrinkling agents, antiseptics; analgesics; antitussives; antipruritics; local anesthetics; anti-hair loss agents; hair growth promoting agents; hair growth inhibitor agents; antihistamines; antiinfectives; inflammation inhibitors; anti-emetics; anticholinergics; vasoconstrictors; vasodilators; wound healing promoters; peptides, polypeptides and proteins; deodorants and anti-perspirants; medicament agents; skin emollients and skin moisturizers; skin firming agents, hair conditioners; hair softeners; hair moisturizers; vitamins; tanning agents; skin lightening agents; antifungals; depilating agents; shaving preparations; external analgesics; perfumes; counterirritants; hemorrhoidals; insecticides; poison ivy products; poison oak products; burn products; anti-diaper rash agents; prickly heat agents; make-up preparations; vitamins; amino acids and their derivatives; herbal extracts; retinoids; flavenoids; sensates; anti-oxidants; skin conditioners; hair lighteners;

chelating agents; cell turnover enhancers; coloring agents; pigments; sunscreens and mixtures thereof.

36. The system of claim 34 wherein the benefit agent is selected from the group consisting of feverfew, centella asiatica, olive leaf, wheat protein, oat oil, lycopene, DMAE, soy and derivatives thereof, colloidal oatmeal, sulfonated shale oil, elubiol, 6-(1-piperidinyl)-2,4-pyrimidinediamine-3-oxide, finasteride, ketoconazole, salicylic acid, zinc pyrithione, coal tar, benzoyl peroxide, selenium sulfide, hydrocortisone, sulfur, menthol, pramoxine hydrochloride, tricetylammonium chloride, polyquaternium 10, panthenol, panthenol triacetate, vitamin A and derivatives thereof, vitamin B and derivatives thereof, vitamin C and derivatives thereof, vitamin D and derivatives thereof, vitamin E and derivatives thereof, vitamin K and derivatives thereof, keratin, lysine, arginine, hydrolyzed wheat proteins, hydrolyzed silk proteins, octyl methoxycinnamate, oxybenzone, minoxidil, titanium dioxide, zinc dioxide, retinol, erthromycin, tretinoin, and mixtures thereof.

37. The system of claim 33 wherein the benefit agent is present in an amount, based upon the total weight of the system, from about 0.001 percent to about 20 percent.

38. The system of claim 25 further comprised of, based upon the total weight of the system, from about 5 percent to about 15 percent of a foaming surfactant.

39. A method of treating hair loss comprising topically applying the system of claim 25 with an effective amount of a hair loss treatment agent to a desired location on an animal or human.

40. The method of claim 39 wherein the hair loss treatment agent is selected from the group consisting of minoxidil, 6-(1-piperidinyl)-2,4-pyrimidinediamine-3-oxide, N'-cyano-N-(tert-pentyl)-N'-3-pyridinyl-guanidine, finasteride, retinoids and derivatives thereof, ketoconazole, elubiol or mixtures thereof.

41. A method for inhibiting hair growth comprising topically applying the system of claim 25 with a hair growth inhibiting agent to a desired area on an animal or human for inhibiting hair growth.

42. The method of claim 41 wherein the benefit agent is selected from the group consisting of serine proteases, retinol, isotretinoin, betamethoisone, alpha-tocophenol and derivatives thereof, and mixtures thereof.

43. A method for treating acne comprising topically applying a mixture of the system of claim 25 and an effective amount of an anti-acne agent to the skin of an animal or human at a desired area.

44. The method of claim 43 wherein the anti-acne agent is selected from the group consisting of benzoyl peroxide, retinol, elubiol, antibiotics, salicylic acid, and mixtures thereof.

45. A method for reducing the signs of aging and other manifestations of photodamage comprising topically applying a mixture of the system of claim 25 and an effective amount of an anti-aging agent to the skin of an animal or human at a desired area.

5 46. The method of claim 45 wherein the anti-aging agent is selected from the group consisting of retinoids, anti-oxidants, alpha-hydroxy acids, beta-hydroxy acids and mixtures thereof.

47. A method for depigmenting the skin comprising topically applying the system of claim 25 and an effective amount of a depigmentation benefit agent to the skin of an animal or human at a desired area.

10 48. The method of claim 47 wherein the depigmentation agent is selected from the group consisting of retinol, Kojic acid, hydroquinone, and mixtures thereof.

49. A method for treating the symptoms and/or the diseases of dandruff, seborrheic dermatitis and/or psoriasis, comprising topically applying a mixture of the system of claim 25 and an effective amount of a benefit agent capable of treating the symptoms to the skin of an animal or human at a desired area.

15 50. The method of claim 49 wherein the benefit agent is selected from the group consisting of shale oil and derivatives thereof, elubiol, ketoconazole, coal tar, salicylic acid, zinc pyrithione, selenium sulfide, hydrocortisone, sulfur, menthol, pramoxine hydrochloride, and mixtures thereof.

20 51. The use of the composition of claim 25 in a personal care product.

52. The composition of claim 51, wherein the personal care product is in the form of a gel, a bath, a wash, a mousse, a shampoo, a rinse, a lotion, a cream, a wipe, a brush, a sponge, or a spray.

53. The use of the composition of claim 25 as a make-up remover.

25 54. The composition of claim 25 in the form of an oil-in-water emulsion.

55. A foaming composition comprising:

a. a water dispersible component;

b. an ester;

c. water; and a

30 d. foaming surfactant.

56. The composition of claim 55 wherein the foaming surfactant has a column height of greater than about 20 mm as determined by the Miles-Ross Test and is selected from the group consisting of non-ionic surfactants, cationic surfactants, amphoteric surfactants, anionic surfactant, and mixtures thereof.

57. The composition of claim 56 further comprising a liquid silicone.

58. The composition of claim 56 further comprising at least one of the following:

- a) polymeric emulsifier and/or a thickener;
- b) a benefit agent; or
- c) a nonionic emulsifier.

5

59. A method for making an oil-in water emulsion comprised of:

a) neutralizing a hydrophilic thickening agent in a hydrophilic phase comprised of a polymeric emulsifier with an effective amount of a neutralizer under conditions sufficient after a lipophilic phase was combined with the hydrophilic phase.

10

60. The method of claim 59 wherein the hydrophilic thickening agent is selected from the group consisting of carbomers, acrylate copolymers, modified hydroxycellulose, polyvinylacetate/maleic anhydride (PVA/MA) decadiene crosspolymer, and mixtures thereof.

15

61. The method of claim 60 wherein the acrylate copolymers are selected from the group consisting of acrylates/aminoacrylates copolymer, acrylates/steareth-20 itaconate copolymer, acrylates/cetheth-20 itaconate copolymer, acrylates/steareth-20 methacrylate copolymer, and mixtures thereof.

62. The method of claim 59 wherein the hydrophilic phase is comprised of one or more of the following components: water, thickener, cleansing enhancer, nonfoaming surfactant, or water dispersible component.

20

63. The method of claim 59 wherein the lipophilic phase is comprised of one or more of the following components: silicone, ester, or polymeric emulsifier.

64. A method for making a water-in-oil emulsion comprised of:

a) neutralizing a hydrophilic thickening agent in a hydrophilic phase comprised of a polymeric emulsifier with an effective amount of a neutralizer under conditions sufficient before combining a lipophilic phase with the hydrophilic phase.

25

65. The method of claim 64 wherein the hydrophilic thickening agent is selected from the group consisting of carbomers, acrylate copolymers, modified hydroxycellulose, polyvinyl methyl ether/maleic anhydride (PVM/MA) decadiene crosspolymer, and mixtures thereof.

30

66. The method of claim 64 wherein the acrylate copolymers are selected from the group consisting of acrylates/aminoacrylates copolymer, acrylates/steareth-20 itaconate

copolymer, acrylates/ceteth-20 itaconate copolymer, acrylates/steareth-20 methacrylate copolymer, and mixtures thereof.

67. The method of claim 64 wherein the hydrophilic phase is comprised of one or more of the following: water, thickener, cleansing enhancer, nonfoaming surfactant, and
5 water dispersible component.

68. The method of claim 64 wherein the lipophilic phase is comprised of one or more of the following: silicone, ester, and polymeric emulsifier.

69. A method for depositing a benefit agent into and/or onto the skin, hair and/or nails comprising applying a composition comprising:

- 10 a. an optional liquid silicone;
- b. a water dispersible component;
- c. an ester;
- d. a polymeric emulsifier and/or a thickener; and
- e. an effective amount of a benefit agent

15 to a desired location on a human or animal.

70. The method of claim 69 wherein the composition is further comprised of a foaming surfactant, a cleansing enhancer, or a mixture thereof.

71. The method of claim 69 wherein the benefit agent is selected from the group consisting of colloidal oatmeal, olive leaf, soy and derivatives thereof, sulfonated shale oil, elubiol, 6-(1-piperidiny)-2,4-pyrimidinediamine-3-oxide, finasteride, ketoconazole, salicylic
20 acid, zinc pyrithione, coal tar, benzoyl peroxide, selenium sulfide, hydrocortisone, sulfur, menthol, pramoxine hydrochloride, tricetylammonium chloride, polyquaternium 10, panthenol, panthenol triacetate, vitamin A and derivatives thereof, vitamin B and derivatives thereof, vitamin C and derivatives thereof, vitamin D and derivatives thereof, vitamin E and
25 derivatives thereof, vitamin K and derivatives thereof, keratin, lysine, arginine, hydrolyzed wheat proteins, hydrolyzed silk proteins, octyl methoxycinnamate, oxybenzone, minoxidil, titanium dioxide, zinc dioxide, retinol, erthromycin, tretinoin, DMAE, and mixtures thereof.

72. A method of cleansing hair or skin comprised of applying an effective amount of the composition of claim 1 to a desired location.

30 73. A method of cleansing hair or skin comprised of applying an effective amount of the system of claim 25 to a desired location.

74. The cleansing composition of claim 25 wherein the water dispersible component is comprised of, based upon the total weight percent of the cleaning system,

- a) from about 0.1 percent to about 5 percent of hexylene glycol;
- b) from about 0.5 percent to about 3.0 percent of polyoxyethylene-6 caprylic/capric triglyceride.

75. A method for depositing a benefit agent into and/or onto the skin, hair and/or
5 nails comprising applying a composition comprising:

- a) an optional liquid silicone;
- b) a water dispersible component
- c) an ester;
- d) water;
- 10 e) a foaming surfactant; and
- f) an effective amount of a benefit agent

to a desired location on a human or animal.

76. A cleaning composition comprised of

- a) a cyclomethicone liquid silicone;
- 15 b) a water dispersible component selected from the group consisting of hexylene glycol, PEG-6 caprylic/capric triglycerides, and mixtures thereof; and
- c) an ester selected from the group consisting of isononyl isononanoate, isostearyl palmitate, cetyl octanoate, pentaerthritol tetraoctanoate, and mixtures thereof.

77. A cleaning system comprised of

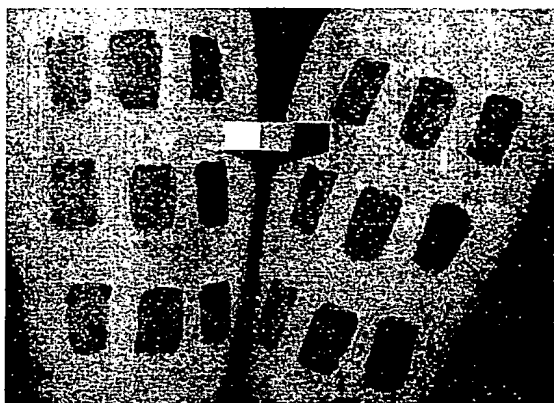
- 20 a) a cyclomethicone liquid silicone;
- b) a water dispersible component selected from the group consisting of hexylene glycol, PEG-6 caprylic/capric triglycerides, and mixtures thereof;
- c) an ester selected from the group consisting of isononyl isononanoate, isostearyl palmitate, cetyl octanoate, pentaerthritol tetraoctanoate, and mixtures thereof;
- 25 d) water; and
- e) a PEG-30 dipolyhydroxystearate polymeric emulsifier and/or a carbomer thickener.

78. A foaming composition comprised of

- a) water;

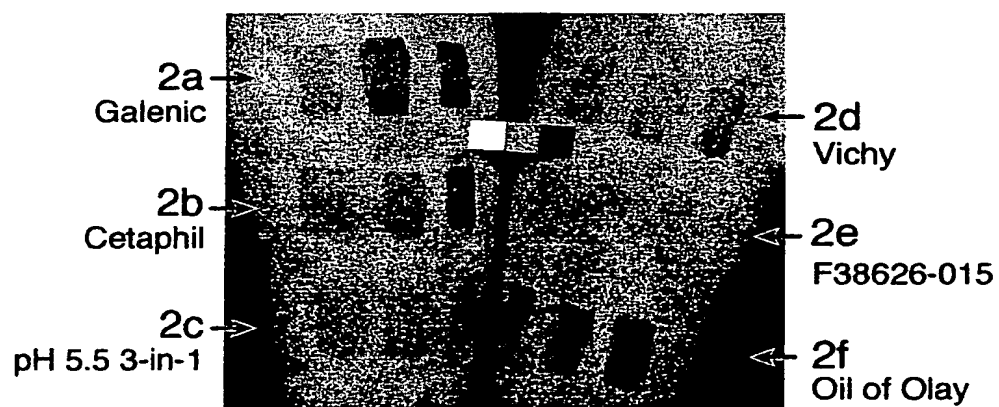
- b) a water dispersible component selected from the group consisting of hexylene glycol, PEG-6 caprylic/capric triglycerides, and mixtures thereof;
- c) an ester selected from the group consisting of isononyl isononanoate, isostearyl palmitate, cetyl octanoate, pentaerthritol tetraoctanoate, and mixtures thereof; and
- 5 d) a foaming surfactant selected from the group consisting of cocamide MEA, lauryl glucoside, PEG-50 tallow amide, cocamodopropylamine oxide, and mixtures thereof.

FIG. 1



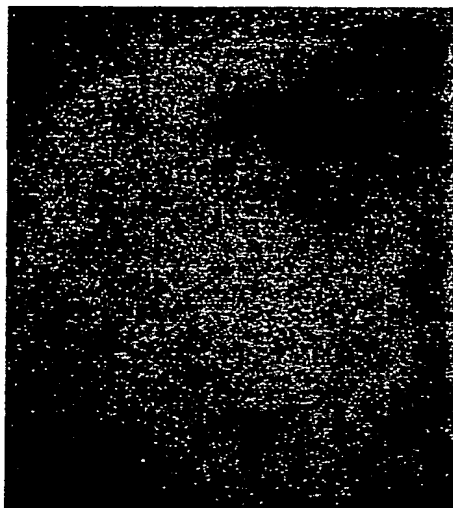
Before Removal

FIG. 2



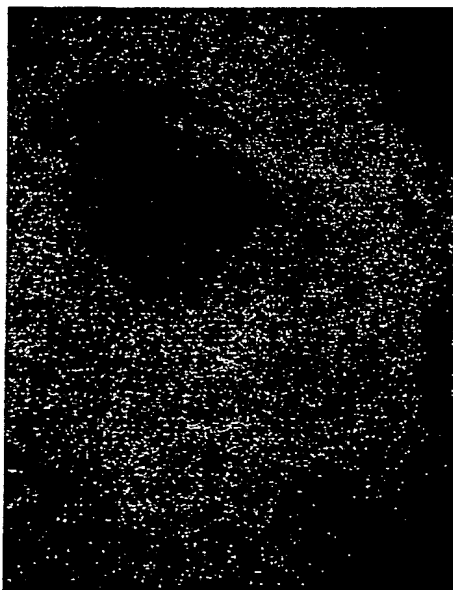
Retinol Fluorescence 0.3% rol Cleanser

FIG. 3(a)



Retinol Fluorescence 0.3% rol Cleanser

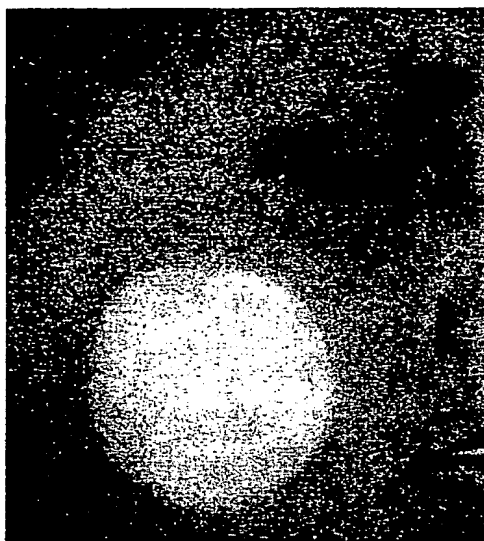
FIG. 3(b)



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Luminosity Change = 78.84

FIG. 3(c)



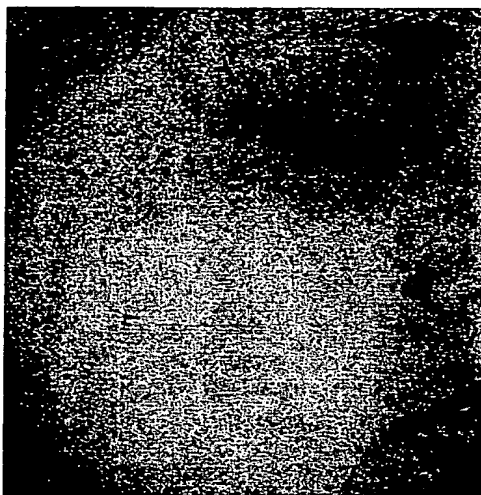
Luminosity Change = 82.33

FIG. 3(d)



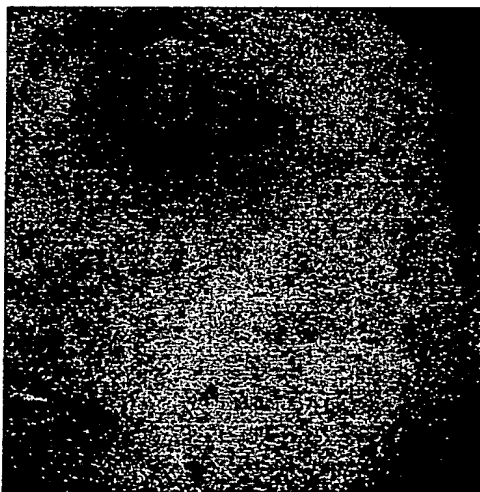
Rinse off Luminosity Change =37.86

FIG. 3(e)



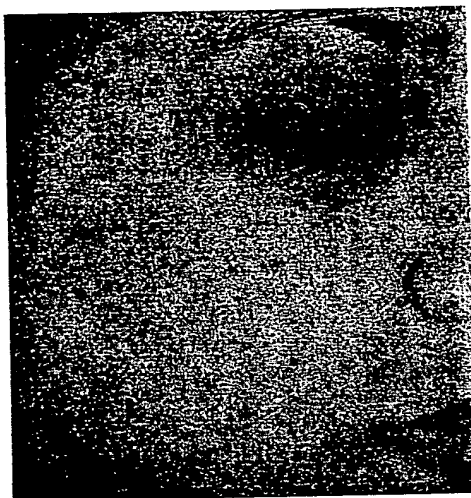
Wiped off Luminosity Change =27.19

FIG. 3(f)



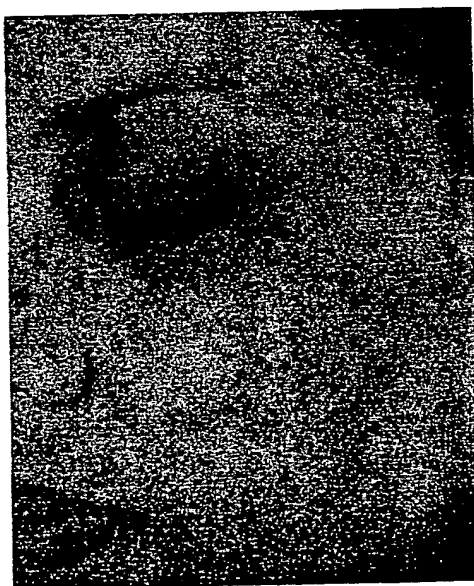
Retinol Fluorescence 0.3% rol Cleanser

FIG. 4(a)



Retinol Fluorescence 0.3% rol Cleanser

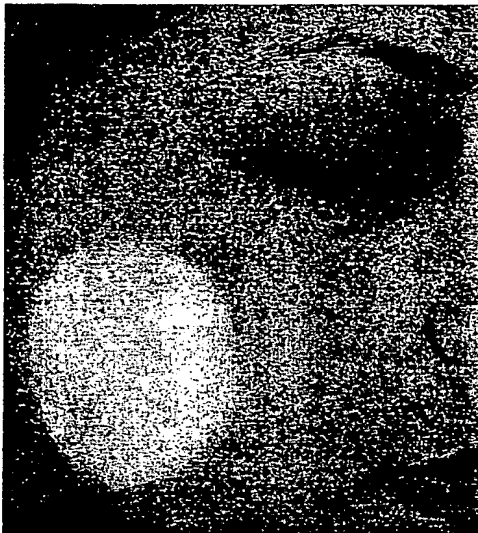
FIG. 4(b)



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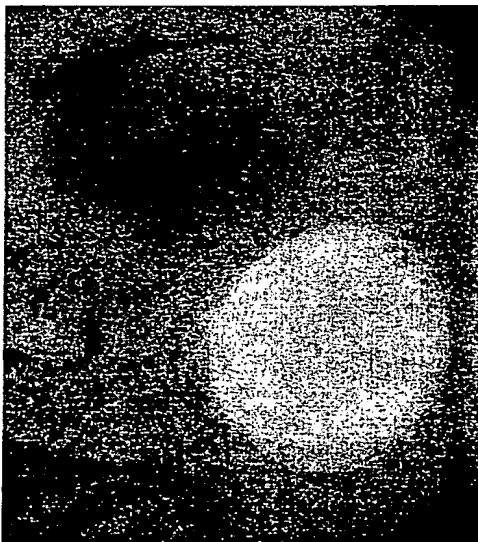
Luminosity Change = 65.92

FIG. 4(c)



Luminosity Change = 70.42

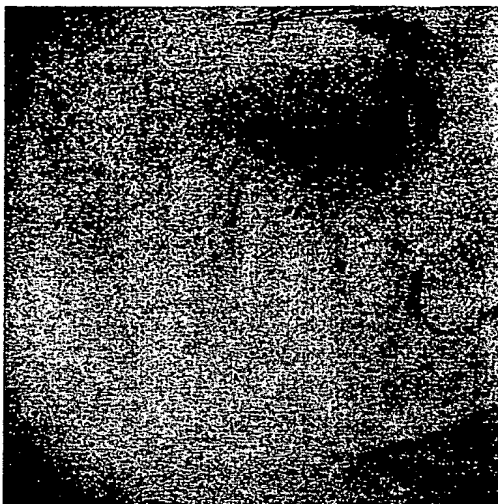
FIG. 4(d)



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Rinse off Luminosity Change =28.97

FIG. 4(e)



Wiped off Luminosity Change =17.01

FIG. 4(f)

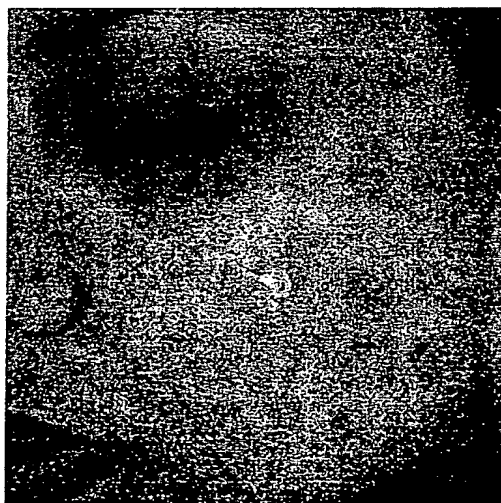


FIG. 5(a)

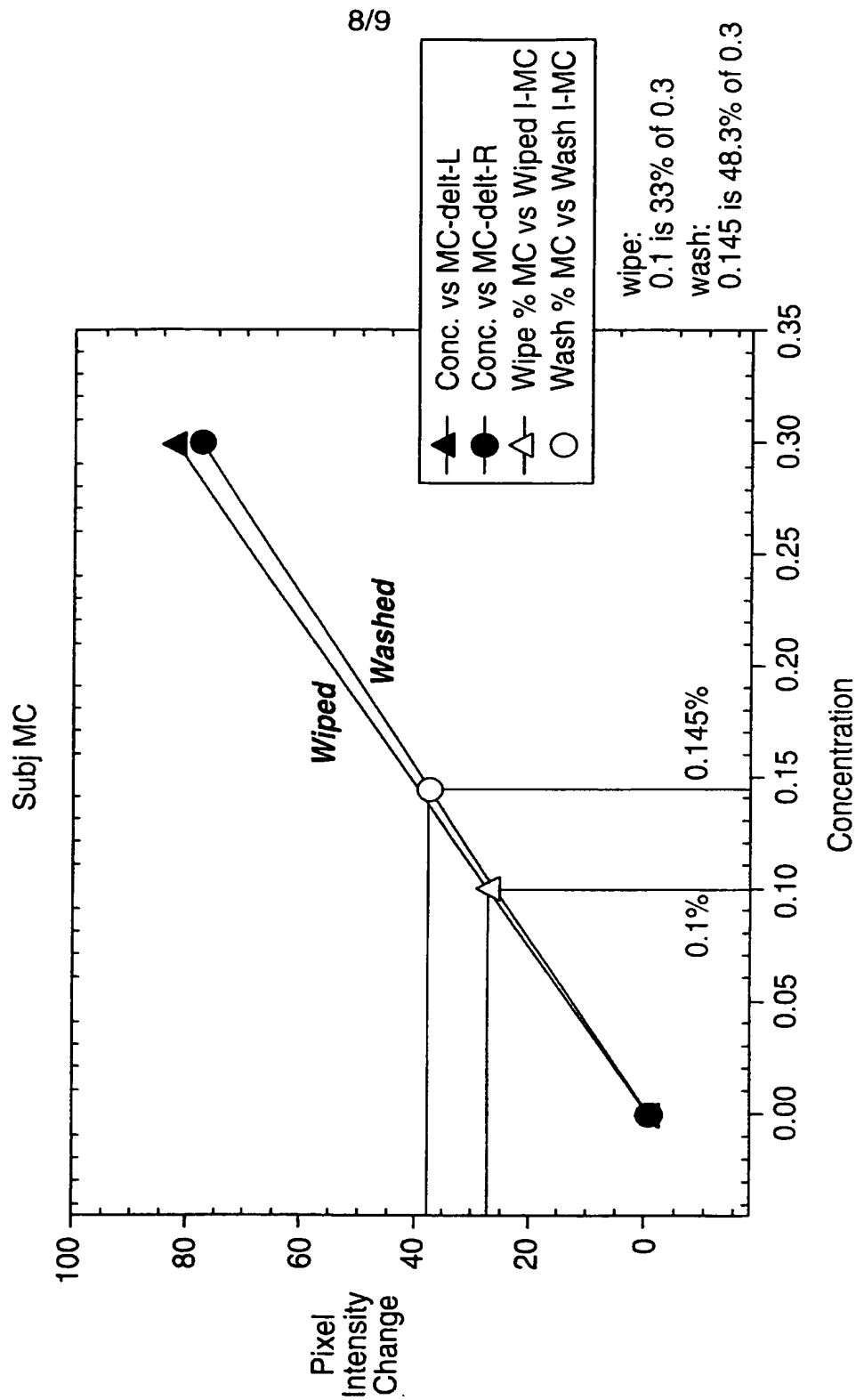
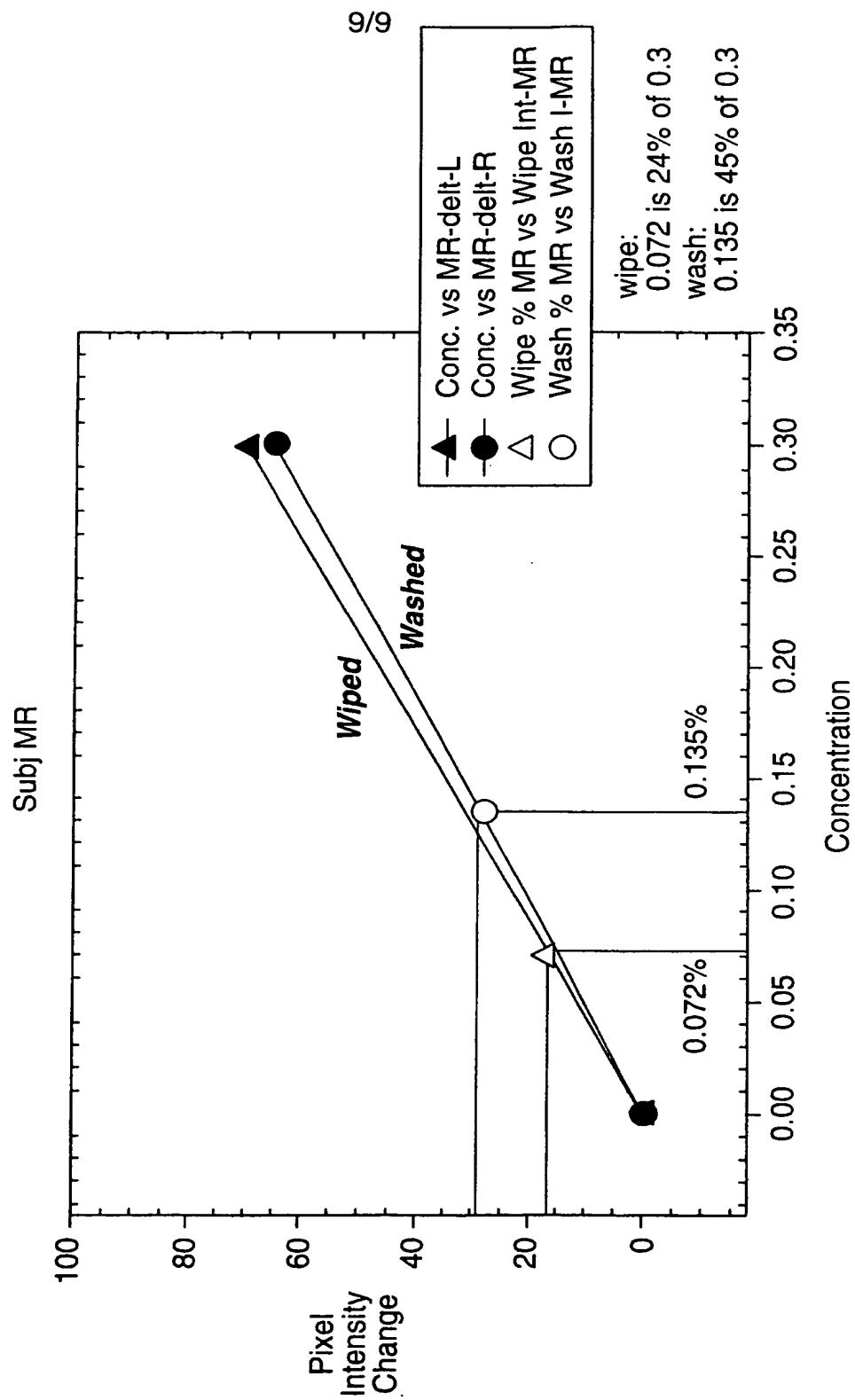


FIG. 5(b)



INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 00/17431

A. CLASSIFICATION OF SUBJECT MATTER IPC 7 A61K7/48 A61K7/50 A61K7/02		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC 7 A61K		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used) WPI Data, PAJ, EPO-Internal		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 95 05160 A (UNILEVER PLC ; UNILEVER NV (NL)) 23 February 1995 (1995-02-23) claims 1,5; example 8 ---	1-26, 55-78
A	WO 98 35652 A (EICKEN ULRICH ; JUNGO SYBILLE (CH); WELLA AG (DE); KISCHKA KARL HEI) 20 August 1998 (1998-08-20) claims 1-3,8-10 ---	1-33, 70-78
A	US 5 888 951 A (SIMON PASCAL ET AL) 30 March 1999 (1999-03-30) claim 1; example 2 ---	1-27, 55-58
A	WO 95 05145 A (UNILEVER PLC ; UNILEVER NV (NL)) 23 February 1995 (1995-02-23) page 9, line 37; claims 1,6 -----	1-26, 51-58, 72,73
<input type="checkbox"/> Further documents are listed in the continuation of box C. <input checked="" type="checkbox"/> Patent family members are listed in annex.		
* Special categories of cited documents : <div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 45%;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"&" document member of the same patent family</p> </div> </div>		
Date of the actual completion of the international search <div style="text-align: center; font-weight: bold;">6 November 2000</div>		Date of mailing of the international search report <div style="text-align: center; font-weight: bold;">17/11/2000</div>
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016		Authorized officer <div style="text-align: center; font-weight: bold;">Beyss, E</div>

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 1,2 (partially)

In present claims 1 and 2 mentioned expressions "water dispersible compound" and "ester" are too broad and relate to an extremely large number of possible products. Support within the meaning of Article 6 PCT and/or disclosure within the meaning of Article 5 PCT is to be found, however, for only a very small proportion of the products claimed. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search over the whole of the claimed scope is impossible. Consequently, the search has been carried out for those parts of the claims which appear to be supported and disclosed, namely those parts relating to the "water dispersible compound" disclosed in claims 5-7 and to the "ester" disclosed in claims 8-24. The general idea of claims 1 and 2 has also been searched.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US. 00/17431

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9505160 A	23-02-1995	AU 677705 B	01-05-1997
		AU 7499294 A	14-03-1995
		AU 7612294 A	14-03-1995
		BR 9407351 A	08-10-1996
		CA 2166468 A	23-02-1995
		DE 69417424 D	29-04-1999
		DE 69417424 T	15-07-1999
		WO 9505159 A	23-02-1995
		EP 0713385 A	29-05-1996
		ES 2129135 T	01-06-1999
		JP 2919973 B	19-07-1999
		JP 9501668 T	18-02-1997
		ZA 9406132 A	15-02-1996
		AU 696398 B	10-09-1998
		AU 7612194 A	14-03-1995
		BR 9407231 A	24-09-1996
		CA 2166469 A	23-02-1995
		WO 9505153 A	23-02-1995
		EP 0793481 A	10-09-1997
		JP 3020278 B	15-03-2000
		JP 9501665 T	18-02-1997
		ZA 9406128 A	17-07-1995
WO 9835652 A	20-08-1998	DE 19705822 A	27-08-1998
		DE 19724587 A	24-12-1998
		BR 9805982 A	31-08-1999
		EP 0897299 A	24-02-1999
US 5888951 A	30-03-1999	FR 2755369 A	07-05-1998
		DE 69700093 D	18-02-1999
		DE 69700093 T	27-05-1999
		EP 0839521 A	06-05-1998
		ES 2129277 T	01-06-1999
		JP 2986436 B	06-12-1999
		JP 10139626 A	26-05-1998
WO 9505145 A	23-02-1995	AU 677704 B	01-05-1997
		AU 7499194 A	14-03-1995
		BR 9407229 A	24-09-1996
		CA 2166532 A	23-02-1995
		DE 69408148 D	26-02-1998
		DE 69408148 T	07-05-1998
		EP 0713381 A	29-05-1996
		ES 2111323 T	01-03-1998
		JP 2944757 B	06-09-1999
		JP 9501667 T	18-02-1997
		ZA 9406131 A	15-02-1996

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(71) Applicant: JOHNSON AND JOHNSON CONSUMER
COMPANIES, INC. [US/US]; 199 Grandview Road,
Skillman, NJ 08558 (US).

(72) Inventors: LUKENBACH, Elvin, R.; 160 Klinesville
Road, Flemington, NJ 08822 (US). KAMINSKI, Claudia;
490 Shire Road, Milford, NJ 08848 (US). PASCAL-SU-
ISSE, Sandrine; Residence Les Annonciades, 5 bis, rue
du Mont, F-76000 Rouen (FR). TAHAR, Maurice; 72,
rue de la Marne, F-27200 Vernon (FR).

(74) Agents: CIAMPORCERO, Audley, A. et al.; Johnson
and Johnson, One Johnson and Johnson Plaza, New
Brunswick, NJ 08933 (US).

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HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,
NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM,
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patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE,
IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG,
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Published:

— With international search report.

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see PCT Gazette No. 12/2001 of 22 March 2001, Section
II

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

(54) Title: CLEANSING COMPOSITIONS

(57) Abstract: Cleansing compositions suitable for use in personal cleansing applications, and in particular make-up removal ap-
plications, which not only impart superior cleansing properties, but also which are relatively non-irritating and thus suitable for use
by people having sensitive skin and eyes comprised of esters, liquid silicones, and a water dispersible components. Also disclosed
are compositions for effectively depositing various benefit agents into and onto the skin.

WO 01/01949 A1

Cleansing Compositions

BACKGROUND OF THE INVENTION

5 1. Field of the Invention

10 This invention relates to cleansing compositions suitable for use in personal cleansing applications, and in particular make-up removal applications, which not only impart superior cleansing properties, but also which are relatively non-irritating and thus suitable for use by people having sensitive skin and eyes. This invention is further related to a composition for effectively delivering and/or depositing various benefit agents into and onto the skin.

2. Description of the Prior Art

15 Various types of cosmetics such as make-up, e.g. lipstick, mascara, foundation, and the like, leave an oil-containing residue on the skin surface that cannot be removed easily by facial cleansers containing conventional soaps. One reason is that such soaps are unable to effectively emulsify or solubilize such oils, which is why many make-up remover compositions have included an oil base as a major component. For example, European Patent No. 370856 discloses a non-foaming makeup remover system comprised of a surfactant-containing water phase that remains physically separated from a cosmetic oil-containing oil phase
20 unless shaken. Disadvantageously, such oil-containing removers also suffer from a tendency to deposit an oily residue or film on the user's skin.

Various attempts have been made to produce stable, oil-free makeup removers. For example, United States Patent No. 5,217,641 discloses an oil-free, stable, non-irritating, single-phase makeup remover comprised of 50 percent to 98 percent of cyclomethicone along with a
25 mixture of esters. However, not only is it economically disadvantageous to use such a large amount of cyclomethicones, but because of their highly volatile nature, cyclomethicones cannot be packaged easily using conventional cosmetic packaging.

Another reason that make-up cannot be removed by conventional soaps is the fact that such soaps are incapable of removing the binders in the make-up. These binders tend to
30 increase the make-up's resistance against sebum and water as well as its overall adhesiveness to the skin. In addition, various polymers, which are similarly difficult to remove, are also employed in hair cosmetics for the purpose of protecting the hair or providing the hair with body.

It would be desirable to have a stable, economically-feasible composition that could not
35 only effectively remove the residue from sebum as well as the residue from make-up and hair-

protecting agents, but also impart a pleasant, non-oily "after-feel" to the skin and hair. It would also be desirable to create such a composition having a low degree of ocular and skin irritation. It would further be desirable to create such a composition that is capable of depositing various active agents into and onto the skin.

5 SUMMARY OF THE INVENTION

In accordance with this invention, there is provided a cleaning composition comprising:

- a. a liquid silicone;
- b. a water dispersible component; and
- c. an ester.

10 Another embodiment of this invention is directed to a cleaning system comprising:

- a. a liquid silicone;
- b. a water dispersible component;
- c. an ester;
- d. water; and
- 15 e. a polymeric emulsifier and/or thickener.

Another embodiment of this invention is directed to a foaming composition comprising:

- a. a water dispersible component;
- b. an ester;
- c. water; and a
- 20 d. foaming surfactant.

Yet another embodiment of the present invention is directed to a method for making an oil-in water emulsion comprised of:

neutralizing a hydrophilic thickening agent in a hydrophilic phase comprised of a polymeric emulsifier with an effective amount of a neutralizer under conditions sufficient after
25 a lipophilic phase was combined with the hydrophilic phase.

Yet another embodiment of the present invention is directed to a method for making a water-in-oil emulsion comprised of:

neutralizing a hydrophilic thickening agent in a hydrophilic phase comprised of a polymeric emulsifier with an effective amount of a neutralizer under conditions sufficient
30 before combining a lipophilic phase with the hydrophilic phase.

Yet another embodiment of the present invention is directed to a method for depositing benefit agents into and onto the skin comprised of:

topically applying an effective amount of the benefit agent with a composition comprised of an optional liquid silicone, a water dispersible component, and an ester to a
5 desired location.

Yet another embodiment of the present invention is directed to a method for depositing a benefit agent into and/or onto the skin, hair and/or nails comprising applying a composition comprising:

- a. an optional liquid silicone;
 - 10 b. a water dispersible component;
 - c. an ester;
 - d. a polymeric emulsifier and/or thickener; and
 - e. an effective amount of a benefit agent
- to a desired location on a human or animal.

15 Yet another embodiment of the present invention is directed to a method for depositing a benefit agent into and/or onto the skin, hair and/or nails comprising applying a composition comprising:

- a. an optional liquid silicone;
- b. a water dispersible component
- 20 c. an ester;
- d. water;
- e. a foaming surfactant; and
- f. an effective amount of a benefit agent

to a desired location on a human or animal.

25 The compositions of this invention are capable of effectively cleansing the skin without significantly contributing to ocular irritation as well as depositing various benefit agents into and onto the skin, hair and nails.

DESCRIPTION OF THE DRAWINGS

30 The file of this patent contains at least one drawing executed in color. Copies of this patent with color drawing(s) will be provided by the Patent and Trademark Office upon request and payment of the necessary fee.

The invention will be more fully understood and further advantages will become apparent when reference is made to the following detailed description of the invention and the accompanying drawing in which:

FIG. 1 is a representation that illustrates two forearms, each of which contains three
5 sets of three different types of makeup before a cleanser is applied thereto.

FIGS. 2 (a) through (f) are representations of the same forearm after six respective cleansers were applied to each set of makeup, respectively.

FIGS. 3 (a) and (b) are representations that illustrate the right side (FIG 3(a)) and left side (FIG. 3(b)) of a subject's face prior to treatment as viewed under a CG-395 Filter.

10 FIGS. 3(c) and (d) are representations that illustrate the right side (FIG 3(c)) and left side (FIG. 3(d)) of a subject's face while possessing the formulation of Example 10 as viewed under a CG-395 Filter.

FIG. 3(e) is a representation that illustrates the right side of a subject's face after the treatment of Example 10 was rinsed therefrom as viewed under a CG-395 Filter.

15 FIG. 3(f) is a representation that illustrates the left side of a subject's face after the treatment of Example 10 was wiped therefrom as viewed under a CG-395 Filter.

FIGS. 4 (a) and (b) are representations that illustrate the right side (FIG 4(a)) and left side (FIG. 4(b)) of a subject's face prior to treatment as viewed under a CG-395 Filter.

20 FIGS. 4(c) and (d) are representations that illustrate the right side (FIG 4(c)) and left side (FIG. 4(d)) of a subject's face while possessing the formulation of Example 10 as viewed under a CG-395 Filter.

FIG. 4(e) is a representation that illustrates the right side of a subject's face after the treatment of Example 10 was rinsed therefrom as viewed under a CG-395 Filter.

25 FIG. 4(f) is a representation that illustrates the left side of a subject's face after the treatment of Example 10 was wiped therefrom as viewed under a CG-395 Filter.

FIG. 5(a) is a graph of concentration of retinol in the formulation of Example 10 versus pixel intensity change.

FIG. 5(b) is a graph of concentration of retinol in the formulation of Example 11 versus pixel intensity change.

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DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

In one embodiment of the present invention, the cleaning composition may suitably comprise, consist of, or consist essentially of, based upon the total weight of the cleaning composition, a) from about 10 percent to about 35 percent, and preferably from about 10 percent to about 20 percent of a liquid silicone; b) from about 10 percent to about 35 percent, and preferably from about 10 percent to about 20 percent of a water dispersible component; and c) from about 30 percent to about 80 percent, and preferably from 55 percent to about 65 percent of an ester.

The first component of the cleaning composition of the present invention may be either a volatile or nonvolatile liquid silicone, with the former being preferred. Examples of suitable silicones nonexclusively include the polydimethyl siloxanes and derivatives thereof such as hexamethylsiloxane, dimethicone, dimethiconol, and cyclomethicone, with cyclomethicone being preferred. Examples of suitable cyclomethicones nonexclusively include cyclotetradimethyl siloxane; cyclopentadimethyl siloxane, cyclohexadimethyl siloxane, cycloheptadimethyl siloxane, and mixtures thereof. Preferably, the silicone has a viscosity of from about 0.25 cp to about 350 cp.

The second component of the cleaning composition of the present invention is a water dispersible component, which is preferably a water soluble solvent. As used herein, the term "water dispersible component" shall mean a material that produces a uniform, clear or hazy, mixture when combined with at least a weight equivalent of water. Examples of suitable water dispersible components nonexclusively include polyethylene glycol 400, hexylene glycol, propylene glycol, polypropylene glycol-10 methylglucose ether, ethoxydiglycol, polyethylene glycol-6 caprylic/capric glyceride, ethylene glycol monobutyl ether, polyethylene glycol-8 caprylic/capric glycerides, 3-methoxy-3-methyl-1-butanol, dimethyl isosorbide, and mixtures thereof. Most preferred water dispersible components include hexylene glycol, dimethyl isosorbide, polyethylene glycol-6 caprylic/capric glyceride, and mixtures thereof.

The third component of the cleaning composition of the present invention is a lipophilic component that preferably is a liquid ester. Preferred esters for use in the composition of this invention include those liquid esters that either possess a structural means for ensuring its liquidity or are heterogeneous in nature. Examples of such structural means include the presence of "interruptions", such as: 1) chain branching; 2) olefinic unsaturation; 3) the presence of either a polyether or a monoalkoxylate in the structure; or 4) the presence of a substituent, e.g. an ethoxylated or propoxylated moiety, bonded between the acid group and the alcohol group. By "heterogeneity," it is meant that the lipophilic component is comprised of a mixture of compounds that vary in the number of carbon atoms in their respective chains.

Examples of suitable esters nonexclusively include:

- a) a branched C₅ to C₂₂ alkyl alcohol ester of an aromatic acid;
 - b) a straight-chained or branched C₅ to C₂₂ alkyl acid esters of optionally ethoxylated/propoxylated polyols having from about 3 carbon atoms to about 7 carbon atoms;
 - c) branched C₅ to C₂₂ alkyl alcohol esters of branched polyacids;
 - d) branched or straight-chained C₅ to C₂₂ alkyl acid esters of branched and/or unsaturated C₅ to C₂₂ alkyl alcohols;
 - e) branched or unsaturated C₅ to C₂₂ alkyl alcohol esters of an acid selected from the group consisting of adipic acid, succinic acid, maleic acid, sebacic acid, and mixtures thereof;
 - f) polyether interrupted fatty acid esters;
 - g) benzoic acid ester of heterogeneous alcohols having from about 8 carbon atoms to about 22 carbon atoms; and
 - h) mixtures thereof,
- with straight-chained or branched C₅ to C₂₂ alkyl acid esters of optionally ethoxylated/propoxylated polyols, benzoic acid esters of heterogeneous alcohols, and mixtures thereof being particularly preferred.

Suitable branched C₅ to C₂₂ alkyl alcohol esters of an aromatic acid include those wherein the aromatic acid is benzoic acid. Preferably, the alcohol of this ester is either branched or unsaturated, and may be either a primary alcohol or a secondary alcohol with the former being preferred. Optionally, the aromatic acid may be substituted with hydroxy or alkyl groups having from about 1 carbon atom to about 4 carbon atoms. Specific examples of these esters nonexclusively include, butyloctyl salicylate; hexyldecyl benzoate; and butyloctyl benzoate, which are all available from C.P. Hall Co. under the tradename, "HallStar," and mixtures thereof, with a mixture of hexyldecyl benzoate and butyloctyl benzoate being particularly preferred.

Another suitable ester includes a straight-chained or branched C₅ to C₂₂ alkyl acid ester of optionally ethoxylated/propoxylated polyols, wherein the polyols contain from about 3 carbon atoms to about 7 carbon atoms. Preferably, if the polyol creates a branching point, then the acid group may be straight-chained. Suitable acids used to form these esters typically have from about 8 carbon atoms to about 22 carbon atoms, and preferably from about 8 carbon atoms to about 18 carbon atoms, and most preferably from about 8 carbon atoms to about 12 carbon atoms, and are either saturated or unsaturated, with octanoic acid,

capric acid, and mixtures thereof being preferred. Such suitable acids are either straight-chained or branched, and are preferably aliphatic. Suitable polyols used to form these esters typically have from about 3 carbon atoms to about 30 carbon atoms, and preferably from about 3 carbon atoms to about 7 carbon atoms. Examples of such suitable polyols
5 nonexclusively include neopentyl alcohol; polyglycerol, e.g. diglycerol, triglycerol, hexaglycerol, and decaglycerol, wherein the polyglycerol may contain from about 2 to about 10 glycerol groups; glycerin; sorbitan; methyl glucose; trimethylolpropane; and mixtures thereof. Neopentyl alcohol, glycerin, trimethylolpropane, and mixtures thereof are the preferred polyols. Examples of suitable esters nonexclusively include pentaerythritol
10 tetraoctanoate; trimethylolpropane trioctanoate; trioctanoin; pentaerythrityl tetrapelargonate; sorbitan trioleate; caprylic/capric triglyceride; neopentyl alcohol tetraoctanoate, and mixtures thereof, with caprylic/capric triglyceride; pentaerythritol tetraoctanoate; trimethylolpropane trioctanoate; and pentaerythrityl tetrapelargonate being more preferred.

Another suitable ester includes the branched C₅ to C₂₂ alkyl alcohol esters of
15 branched polyacids such as the tri-esters, tetra-esters, penta-esters, and mixtures thereof. An example of such a polyacid is citric acid. Suitable alkyl alcohols for creating these esters are optionally substituted, e.g., ethoxylated or propoxylated, contain from about 3 carbon atoms to about 22 carbon atoms, and preferably from about 3 carbon atoms to about 8 carbon atoms, and are either straight-chained or branched, with the branching being
20 preferred. These alcohols may either be primary or secondary, and may either be saturated or unsaturated, with saturated being preferred for stability reasons. Specific examples of suitable esters nonexclusively include trioctyldodecyl citrate; triisopropylcitrate; and mixtures thereof.

Another suitable ester includes the branched or straight-chained C₅ to C₂₂ alkyl acid
25 esters of branched or unsaturated alkyl alcohols wherein the alkyl group of the alcohol has from about 1 carbon atoms to about 18 carbon atoms, and preferably from about 4 carbon atoms to about 10 carbon atoms, provided that the total number of carbon atoms in the ester is at least about 8. Suitable acids for use in making these esters typically have from about 2 carbon atoms to about 22 carbon atoms, and preferably from about 5 carbon atoms to about
30 10 carbon atoms. However, if the number of acid carbon atoms exceeds the number of carbon atoms in the alcohol, then the acid preferably contains from about 8 carbon atoms to about 18 carbon atoms and the alcohol preferably contains from about 1 carbon atom to about 8 carbon atoms. If the number of acid carbon atoms is less than the number of carbon atoms in the alcohol, then the acid preferably contains from about 2 carbon atoms to about 8
35 carbon atoms and the alcohol preferably contains from about 8 carbon atoms to about 18 carbon atoms. Preferably, either: 1) the alcohol group or the acid group has branching and/or unsaturation, i.e. both the alcohol and the acid are not straight-chained; or 2) the

ester possesses an asymmetrical alkyl distribution. By "asymmetrical alkyl distribution," it is meant that the ester is made from, for example, a short chain alcohol, i.e. having from about 1 carbon atom to about 8 carbon atoms, and a long chain acid, i.e., having greater than about 8 carbon atoms, such as, e.g. butyl stearate, or less preferably the ester is made from, a long chain alcohol, i.e. having greater than about 8 carbon atoms, and a short chain acid, i.e. having from about 1 carbon atom to about 8 carbon atoms. Examples of such suitable esters nonexclusively include tridecyl neopentanoate, isostearyl palmitate, cetyl ricinoleate, cetyl octanoate, isononyl isononanoate, butyl stearate, octyldodecyl soyate, tridecyl erucate, octyldodecyl erucate/eicosil erucate, and mixtures thereof, with cetyl octanoate, isostearyl palmitate, isononyl isononanoate, and mixtures thereof and being preferred.

Another suitable ester includes the branched or unsaturated C₅ to C₂₂ alkyl alcohol esters of an acid selected from the group consisting of adipic acid, succinic acid, maleic acid, sebacic acid, and mixtures thereof. The alcohol of these esters, which has from about 3 carbon atoms to about 18 carbon atoms, and preferably from about 3 carbon atoms to about 8 carbon atoms, is preferably branched or unsaturated. Examples of such suitable alcohol esters nonexclusively include diisopropyl adipate, dioctyl sebacate, dioctyl succinate, dioctyl maleate, diisostearyl adipate, diethyl sebacate, and mixtures thereof, with diethyl sebacate, dioctyl sebacate, and diisostearyl adipate being preferred.

Another suitable ester includes polyether interrupted fatty acid esters. Examples of such suitable esters nonexclusively include: 1) laureth-2 benzoate; 2) C₈ to C₂₂ fatty alkyl (optionally polypropylenoxy) polyethyleneoxy carboxylate esters derived from an alcohol having from about 1 carbon atom to about 22 carbon atoms, is either straight or branched, and may contain a phenyl group; and 3) mixtures thereof, with C₈ to C₂₂ fatty alkyl (optionally polypropylenoxy) polyethyleneoxy carboxylate esters being preferred. Specific examples of preferred esters nonexclusively include isopropyl propylene glycol-2-isodeceth-7 carboxylate, such as "Velsan D8P3" and other commercially available materials sold by Sandoz under the tradename, "Velsan."

Another suitable ester includes the benzoic acid esters of heterogeneous alcohols having from about 8 carbon atoms to about 22 carbon atoms, such as the ester mixtures available from Finetex under the tradename, "Finsolv" and preferably is the C₁₂ to C₁₅ alcohols benzoate available from Finetex under the tradename, "Finsolv TN."

Preferred combinations of esters include at least one, preferably at least two, and more preferably three of the following esters: a) branched C₅ to C₂₂ alkyl alcohol esters of an aromatic acid; b) branched or straight-chained C₅ to C₂₂ alkyl acid esters of branched or unsaturated alkyl alcohols; and c) straight-chained or branched C₅ to C₂₂ alkyl acid esters of optionally ethoxylated/propoxylated polyols. In a preferred embodiment, the ester

contains, based upon the total weight percent of the esters, from about 30 percent to about 80 percent of branched or straight-chained C₅ to C₂₂ alkyl acid esters of branched or unsaturated C₅ to C₂₂ alkyl alcohols; from about 10 percent to about 50 percent of branched C₅ to C₂₂ alkyl alcohol esters of an aromatic acid; and from about 10 percent to about 50 percent of straight-chained or branched C₅ to C₂₂ alkyl acid esters of optionally ethoxylated/propoxylated polyols. In a more preferred embodiment, the ester contains, based upon the total weight percent of the esters, from about 15 percent to about 50 percent isononyl isononanoate, from about 15 percent to about 50 percent isostearyl palmitate, from about 15 percent to about 50 percent cetyl octanoate, and from about 15 percent to about 50 percent pentaerthritol tetraoctanoate.

Another embodiment of the present invention is directed to a cleaning system comprising, consisting, or consisting essentially of, based upon the total weight of the cleaning system, a) at least 5 percent and preferably at least about 10 percent of the cleaning composition; b) from about 70 percent to about 98 percent, and preferably from about 80 percent to about 90 percent of water; c) from about 0.1 percent to 5 percent, e.g. from about 0.5 percent to about 1.5 percent of a polymeric emulsifier, a thickener, or mixture thereof; optionally d) from about 0.1 percent to about 5 percent, and preferably from about 1 percent to about 3 percent of a cleaning enhancer; optionally e) from about 2 percent to about 20 percent, and preferably from about 5 percent to about 15 percent of a foaming surfactant; and optionally f) from about 0.001 percent to about 20 percent of a benefit agent. In one embodiment, the cleaning system may comprise, based upon the total weight of the cleansing system, from about 0.1 to about 5 percent, and preferably from about 0.5 percent to 1.5 percent of a polymeric emulsifier and/or from about 0.01 percent to about 2 percent, and preferably from about 0.01 percent to about 0.5 percent of a thickener. More preferably, the cleaning system contains, based upon the total weight of the cleaning system, from about 10 percent to about 30 percent of the cleaning composition.

The cleaning system may be in the form of an oil-in-water emulsion, a water-in-oil emulsion, or a dispersion.

In addition to the cleansing composition, the cleaning system is further comprised of polymeric emulsifiers and/or thickeners. As used herein, the term "polymeric emulsifier" shall mean those compounds capable of emulsifying cleaning systems whereby the polymeric emulsifiers have a molecular weight of at least about 5000, and preferably are block copolymers having a hydrophilic portion and a hydrophobic portion. When used at amounts effective for emulsifying the cleansing system, the polymeric emulsifiers surprisingly do not cause significant eye sting, i.e., when the emulsifier-containing composition was used by 80 consumers in the eye area, no more than about 5% of such users expressed discomfort

around the eye area. Examples of suitable polymeric emulsifiers nonexclusively include polyethylene glycol-30 dipolyhydroxystearate available from Uniqema under the tradename, 'Arlacel P-135;" dimethicone copolyol, which is available from Goldschmidt Chemical Corporation under the tradename, "Abil EM 90"; substituted acrylates such as those available
5 from The Goodrich Corporation under the tradename, "Pemulen"; and mixtures thereof, with polyethylene glycol-30 dipolyhydroxystearate being preferred.

Examples of suitable hydrophilic thickeners nonexclusively include carbomers available from B.F. Goodrich under the tradename, "Carbopol ETD 2020", acrylate copolymers, hydroxyethylcellulose modified with cetyl ether groups available from Hercules
10 under the tradename, "Natrosol Plus", polyvinylmethyl ether/maleic anhydride (PVM/MA) decadiene crosspolymer available from International Specialty Products under the tradename, "Stabileze QM," and copolymers and mixtures thereof, with carbomers being preferred. Examples of suitable acrylate copolymers nonexclusively include acrylate copolymers available from Rohm & Haas under the tradename, "Aculyn 33,"
15 acrylates/aminoacrylates copolymer available from National Starch & Chemical Company under the tradename, "Structure Plus," acrylates/steareth-20 itaconate copolymer available from National Starch & Chemical Company under the tradename, "Structure 2001," acrylates/ceteth-20 itaconate copolymer available from National Starch & Chemical Company under the tradename, "Structure 3001," acrylates/steareth-20 methacrylate
20 copolymer available from Rohm & Haas under the tradename, "Aculyn 22," and copolymers and mixtures thereof.

The cleaning system of the present invention may also optionally contain a cleaning enhancer in the form of a nonionic emulsifier and/or a non-foaming surfactant. Examples of suitable nonionic emulsifiers include isocetheth-20, oleth-2, mixture of PEG-40 hydrogenated
25 castor oil and trideceth-9 available from Dragoco Inc. under the tradename, "Dragoco Solubilizer 2/014160," Poloxamer 184, laureth-4, sorbitan trioleate, polyoxyethylene-(2) oleyl ether, sorbitan stearate, cetearyl glucoside, glyceryl oleate, trideceth-9, polyethylene glycol-40 hydrogenated castor oil, and mixtures thereof.

Examples of suitable non-foaming surfactants include non-foaming nonionic
30 surfactants such as sucrose esters, e.g., sucrose cocoate, sucrose stearate and mixtures thereof, with sucrose cocoate being preferred. By "essentially non-foaming," it is meant that the surfactant, when used with the composition of the present invention, has a column height of less than about 20 mm as determined by the Ross-Miles Foam Generation Test. See 18 (I.) Oil & Soap 99 – 102 (1941) ["Ross-Miles Test"], which is incorporated by reference
35 herein. The cleansing composition and the cleansing system may either be rinseable with water or may be wiped-off. Preferably, the essentially, non-foaming surfactants are used in

embodiments wherein the cleansing system or the cleansing composition is rinseable with water. For example, a preferred combination of hydrophilic components include, based upon the total weight percent of the cleansing system, from about 0.1 percent to about 5.0 percent of hexylene glycol, from about 0.5 percent to about 3.0 percent of sucrose cocoate non foaming surfactant, and from about 0.5 percent to about 3.0 percent of polyoxyethylene-6 caprylic/capric triglyceride. An example of a suitable cleaning enhancer include a mixture of sorbitan stearate and sucrose cocoate available from Uniqema under the tradename, "Arlatone 2121."

Preferably, the cleaning system contains, based upon the total weight of the cleansing system, no more than about 6%, and preferably 5%, of the cleaning enhancers for cream formulations and no more than about 2%, and preferably no more than 1% of the cleaning enhancers in thin lotion/milk formulations.

The cleansing system and cleansing composition may also optionally contain a foaming surfactant. The foaming surfactant may be non-ionic, cationic, amphoteric, or anionic; nonionic surfactants are preferred. By "foaming," it is meant that the surfactant, when used with the composition of the present invention, has a column height of foam greater than about 20 mm as determined by the Ross-Miles Test. As used herein, the term "amphoteric" shall mean: 1) molecules that contain both acidic and basic sites such as, for example, an amino acid containing both amino (basic) and acid (e.g., carboxylic acid, acidic) functional groups; or 2) zwitterionic molecules which possess both positive and negative charges within the same molecule. The charges of the latter may be either dependent on or independent of the pH of the composition. Examples of zwitterionic materials include, but are not limited to, alkyl betaines and amidoalkyl betaines. Examples of suitable and preferred surfactants may be found in International Patent Application Number WO97/01196, which is incorporated by reference in its entirety herein.

The cleansing system may further contain one or more benefit agents or pharmaceutically-acceptable salts thereof. As used herein, the term "benefit agent" includes any active ingredient that is to be delivered into and/or onto the skin, hair or nail at a desired location, such as a cosmetic agent or a pharmaceutical agent. By "cosmetic agent," it is meant any ingredient that is appropriate for cosmetically treating, providing nutrients to, and/or conditioning the hair, nail, and/or skin via topical application. By "pharmaceutical agent," it is mean any drug that is either hydrophobic or hydrophilic in nature and appropriate for topical use. As used herein "medicament agents" include those agents capable of promoting recovery from injury and illness.

The benefit agents useful herein may be categorized by their therapeutic benefit or their postulated mode of action. However, it is to be understood that the benefit agents

useful herein may, in some circumstances, provide more than one therapeutic benefit or operate via greater than one mode of action. Therefore, the particular classifications provided herein are made for the sake of convenience and are not intended to limit the benefit agents to the particular application(s) listed. In addition, the compounds, which are
5 identified below as being suitable for use as benefit agents, may be used in an amount over and above the amount that they may be used for other purposes in the cleansing composition/cleansing system.

Examples of suitable benefit agents include, but are not limited to, depigmentation agents; reflectants; detangling/wet combing agents; film forming polymers; humectants;
10 amino acids and their derivatives; antimicrobial agents; allergy inhibitors; anti-acne agents; anti-aging agents; anti-wrinkling agents, antiseptics; analgesics; antitussives; antipruritics; local anesthetics; anti-hair loss agents; hair growth promoting agents; hair growth inhibitor agents, antihistamines such as *Mandragora Vernalis*, *Tanacetum Parthenium* and the like; antiinfectives such as *Acacia Catechu*, *Aloe Barbadensis*, *Convallaria Majalis*, *Echinacea*,
15 *Eucalyptus*, *Mentha Piperita*, *Rosa Canina*, *Sassafras Albidum*, and the like; inflammation inhibitors; anti-emetics; anticholinergics; vasoconstrictors; vasodilators; wound healing promoters; peptides, polypeptides and proteins; deodorants and anti-perspirants; medicament agents; skin emollients and skin moisturizers; skin firming agents, hair conditioners; hair softeners; hair moisturizers; vitamins; tanning agents; skin lightening
20 agents; antifungals such as *Centaurea Cyanus*, *Kalmia Latifolia* and antifungals for foot preparations; depilating agents; shaving preparations; external analgesics; perfumes; counterirritants; hemorrhoidals; insecticides; poison ivy products; poison oak products; burn products; anti-diaper rash agents; prickly heat agents; make-up preparations; vitamins; amino acids and their derivatives; herbal extracts; retinoids; flavenoids; sensates; anti-
25 oxidants; skin conditioners; hair lighteners; chelating agents; cell turnover enhancers; coloring agents; pigments; sunscreens, those active ingredients disclosed in United States Patent No. 6,063,397, which is incorporated herein by reference, anti-edema agents, collagen enhancers, and mixtures thereof.

Examples of suitable anti-edema agents nonexclusively include bisabolol natural,
30 synthetic bisabolol, and mixtures thereof.

Examples of suitable vasoconstrictors nonexclusively include horse chestnut extract, prickly ash, and mixtures thereof.

Examples of suitable anti-inflammatory agents nonexclusively include benoxaprofen, centella asiatica, bisabolol, feverfew (whole), feverfew (parthenolide free), green tea extract,
35 green tea concentrate, hydrogen peroxide, lycopene including "Lyc-o-Pen" available from LycoRed Natural Products Industries, Ltd., oat oil, chamomile, and mixtures thereof.

Examples of collagen enhancers nonexclusively include vitamin A, vitamin C, and mixtures thereof.

Examples of suitable skin firming agent nonexclusively include dimethylaminoethanol ("DMAE").

5 Examples of suitable antipruritics and skin protectants nonexclusively include oatmeal, betaglucan, feverfew, soy and derivatives thereof, bicarbonate of soda, colloidal oatmeal, surfactant based colloidal oatmeal cleanser, Anagallis Arvensis, Oenothera Biennis, Verbena Officinalis, and the like. These antipruritics may be used in an amount, based upon the total weight of the cleansing composition, from about 0.01 percent to about
10 40 percent, and preferably from about 1 percent to about 5 percent.

As used herein, colloidal oatmeal means the powder resulting from the grinding and further processing of whole oat grain meeting United States Standards for Number 1 or Number 2 oats. The colloidal oatmeal has a particle size distribution as follows: not more than 3 percent of the total particles exceed 150 micrometers in size and not more than 20
15 percent of the total particles exceed 75 micrometers in size. Examples of suitable colloidal oatmeals include, but are not limited to, "Tech-O" available from the Beacon Corporation and colloidal oatmeals available from Quaker.

Examples of suitable reflectants nonexclusively include mica, alumina, calcium silicate, glycol dioleate, glycol distearate, silica, sodium magnesium fluorosilicate, and
20 mixtures thereof.

Suitable detangling/wet combing agents nonexclusively include polyquaternium-10, hydroxypropyltrimonium guar, dioleoylamidoethyl hydroxyethylmonium methosulfate, di- (soyoylethyl) hydroxyethylmonium methosulfate, hydroxyethyl behenamidopropyl dimonium chloride, olealkonium chloride, polyquaternium-47, stearylalkonium chloride, tricetylmonium
25 chloride, and mixtures thereof.

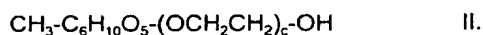
Suitable film forming polymers include those that, upon drying, produce a substantially continuous coating or film on the hair, skin, or nails. Nonexclusive examples of suitable film forming polymers include acrylamidopropyl trimonium chloride/acrylamide copolymer; corn starch/ acrylamide/ sodium acrylate copolymer; polyquaternium-10; polyquaternium-47;
30 polyvinylmethylether/maleic anhydride copolymer; styrene/acrylates copolymers; and mixtures thereof.

Commercially available humectants which are capable of providing moisturization and conditioning properties to the cleansing composition are suitable for use in the present invention. The humectant is preferably present in an amount of from about 0 percent to about
35 10 percent, more preferably from about 0.5 percent to about 5 percent, and most preferably from about 0.5 percent to about 3 percent, based on the overall weight of the composition. Examples of suitable humectants nonexclusively include: 1) water soluble liquid polyols

selected from the group comprising glycerine, propylene glycol, hexylene glycol, butylene glycol, pentylene glycol, dipropylene glycol, and mixtures thereof; 2) polyalkylene glycol of the formula I.:



- 5 wherein R'' is an alkylene group having from about 2 to about 4 carbon atoms and b is an integer of from about 1 to about 10, such as PEG 4; 3) polyethylene glycol ether of methyl glucose of formula II.:



wherein c is an integer from about 5 to about 25;

- 10 4) urea; 5) fructose; 6) glucose; 7) honey; 8) lactic acid; 9) maltose; 10) sodium glucuronate; and 11) mixtures thereof, with glycerine being the preferred humectant.

Suitable amino acid agents include amino acids derived from the hydrolysis of various proteins as well as the salts, esters, and acyl derivatives thereof. Examples of such amino acid agents nonexclusively include amphoteric amino acids such as alkylamido
15 alkylamines, i.e. stearyl acetyl glutamate, capryloyl silk amino acid, capryloyl collagen amino acids; capryloyl keratin amino acids; capryloyl pea amino acids; cocodimonium hydroxypropyl silk amino acids; corn gluten amino acids; cysteine; glutamic acid; glycine; hair keratin amino acids; amino acids such as aspartic acid, threonine, serine, glutamic acid, proline, glycine, alanine, cystine, valine, methionine, isoleucine, leucine, tyrosine,
20 phenylalanine, cysteic acid, lysine, histidine, arginine, cysteine, tryptophan, citrulline; lysine; silk amino acids, wheat amino acids; and mixtures thereof

Suitable proteins include those polymers that have a long chain, i.e. at least about 10 carbon atoms, and a high molecular weight, i.e. at least about 1000, and are formed by self-condensation of amino acids. Nonexclusive examples of such proteins include collagen,
25 deoxyribonuclease, iodized corn protein; milk protein; protease; serum protein; silk; sweet almond protein; wheat germ protein; wheat protein; alpha and beta helix of keratin proteins; hair proteins, such as intermediate filament proteins, high-sulfur proteins, ultrahigh-sulfur proteins, intermediate filament-associated proteins, high-tyrosine proteins, high-glycine tyrosine proteins, trichohyalin, and mixtures thereof.

30 Examples of suitable vitamins nonexclusively include vitamin B complex; including thiamine, nicotinic acid, biotin, pantothenic acid, choline, riboflavin, vitamin B6, vitamin B12, pyridoxine, inositol, carnitine; vitamins A,C,D,E,K and their derivatives such as vitamin A palmitate and pro-vitamins, e.g. (i.e. panthenol (pro vitamin B5) and panthenol triacetate) and mixtures thereof.

Examples of suitable antibacterial agents nonexclusively include bacitracin, erythromycin, neomycin, tetracycline, chlortetracycline, benzethonium chloride, phenol, and mixtures thereof.

5 Examples of suitable skin emollients and skin moisturizers nonexclusively include mineral oil, lanolin, vegetable oils, isostearyl isostearate, glyceryl laurate, methyl gluceth-10, methyl gluceth-20 chitosan, and mixtures thereof.

10 Examples of suitable hair conditioners nonexclusively include quaternized compounds such as behenamidopropyl PG-dimonium chloride, tricetylmonium chloride, dihydrogenated tallowamidoethyl hydroxyethylmonium methosulfate, and mixtures thereof as well as lipophilic compounds like cetyl alcohol, stearyl alcohol, hydrogenated polydecene, and mixtures thereof.

15 An example of a suitable hair softener nonexclusively includes silicone compounds, such as those that are either non-volatile or volatile and those that are water soluble or water insoluble. Examples of suitable silicones include organo-substituted polysiloxanes, which are either linear or cyclic polymers of monomeric silicone/oxygen monomers and which nonexclusively include cetyl dimethicone; cetyl triethylammonium dimethicone copolyol phthalate; cyclomethicone; dimethicone copolyol; dimethicone copolyol lactate; hydrolyzed soy protein/dimethicone copolyol acetate; silicone quaternium 13; stearammonium dimethicone copolyol phthalate; stearamidopropyl dimethicone; and mixtures thereof.

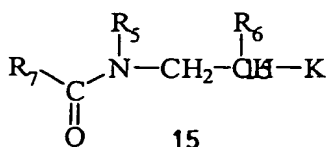
20 Examples of suitable hair moisturizers nonexclusively include panthenyl ethyl ether, phytantriol, and mixtures thereof.

25 Examples of sunscreen agents nonexclusively include benzophenones, borrelone, butyl paba, cinnamidopropyl trimethyl ammonium chloride, disodium distyrylbiphenyl disulfonate, paba, potassium methoxycinnamate, butyl methoxydibenzoylmethane, octyl methoxycinnamate, oxybenzone, octocrylene, octyl salicylate, phenylbenzimidazole sulfonic acid, ethyl hydroxypropyl aminobenzoate, menthyl anthranilate, aminobenzoic acid, cinoxate, diethanolamine methoxycinnamate, glyceryl aminobenzoate, titanium dioxide, zinc oxide, oxybenzone, Padimate O, red petrolatum, and mixtures thereof.

30 An example of a suitable tanning agent nonexclusively includes dihydroxyacetone.

Examples of skin lightening agents nonexclusively include hydroquinone, catechol and its derivatives, ascorbic acid and its derivatives, and mixtures thereof.

35 Examples of suitable insecticides (including insect repellents, anti-scabies and anti-lice treatments) nonexclusively include permethrin, pyrethrin, piperonyl butoxide, imidacloprid, N,N-diethyl toluamide, which refers to the material containing predominantly the *meta* isomer, i.e., N,N-diethyl-*m*-toluamide, which is also known as DEET; compounds of the formula III.



III.

wherein

- 5 R_5 is a branched or unbranched alkyl group having about 1 to about 6 carbon atoms;
- R_6 is H, methyl or ethyl;
- R_7 is a branched or unbranched alkyl or alkoxy group having from about 1 to about 8 carbon atoms; and
- 10 K is a -CN or a $-COOR_8$ group, wherein

R_8 is a branched or unbranched alkyl group having from about 1 to about 6 carbon atoms,

- natural or synthetic pyrethroids, whereby the natural pyrethroids are contained in pyrethrum, the extract of the ground flowers of *Chrysanthemum cinerariaefolium* or *C. coccineum*; and mixtures thereof. Within the structure of Formula III. are ethyl 3-(N-
- 15 butylacetamido)propionate, wherein R_7 is a CH_3 group, R_5 is an n-butyl group, R_6 is H, K is $COOR_8$ and R_8 is ethyl, which is available commercially from Merck KGaA of Darmstadt, Germany under the name, "Insect Repellent 3535."

- An example of an anti fungal for foot preparations nonexclusively includes tolnaftate.
- 20 Examples of suitable depilating agents nonexclusively include calcium thioglycolate, magnesium thioglycolate, potassium thioglycolate, strontium thioglycolate, and mixtures thereof.

- Examples of suitable external analgesics and local anesthetics nonexclusively include benzocaine, dibucaine, benzyl alcohol, camphor, capsaicin, capsicum, capsicum
- 25 oleoresin, juniper tar, menthol, methyl nicotinate, methyl salicylate, phenol, resorcinol, turpentine oil, and mixtures thereof.

 Examples of suitable antiperspirants and deodorants nonexclusively include aluminium chlorohydrates, aluminium zirconium chlorohydrates, and mixtures thereof.

- Examples of suitable counterirritants nonexclusively include camphor, menthol,
- 30 methyl salicylate, peppermint and clove oils, ichtammol, and mixtures thereof.

 An example of a suitable inflammation inhibitor nonexclusively includes hydrocortisone, *Fragaria Vesca*, *Matricaria Chamomilla*, and *Salvia Officinalis*.

 Examples of suitable hemorrhoidal products nonexclusively include the anesthetics such as benzocaine, pramoxine hydrochloride, and mixtures thereof; antiseptics such as

benzethonium chloride; astringents such as zinc oxide, bismuth subgallate, balsam Peru, and mixtures thereof; skin protectants such as cod liver oil, vegetable oil, and mixtures thereof.

Most preferred benefit agents nonexclusively include DMAE, soy and derivatives thereof, colloidal oatmeal, sulfonated shale oil, olive leaf, elubiol, 6-(1-piperidiny)-2,4-pyrimidinediamine-3-oxide, finasteride, ketoconazole, salicylic acid, zinc pyrithione, coal tar, benzoyl peroxide, selenium sulfide, hydrocortisone, sulfur, menthol, pramoxine hydrochloride, tricetylmonium chloride, polyquaternium 10, panthenol, panthenol triacetate, vitamin A and derivatives thereof, vitamin B and derivatives thereof, vitamin C and derivatives thereof, vitamin D and derivatives thereof, vitamin E and derivatives thereof, vitamin K and derivatives thereof, keratin, lysine, arginine, hydrolyzed wheat proteins, hydrolyzed silk proteins, octyl methoxycinnamate, oxybenzone, minoxidil, titanium dioxide, zinc dioxide, retinol, erythromycin, tretinoin, and mixtures thereof.

One preferred type of benefit agent includes those therapeutic components that are effective in the treatment of dandruff, seborrheic dermatitis, and psoriasis as well as the symptoms associated therewith. Examples of such suitable benefit agents nonexclusively include zinc pyrithione, anthralin, shale oil and derivatives thereof such as sulfonated shale oil, selenium sulfide, sulfur, salicylic acid, coal tar, povidone-iodine, imidazoles such as ketoconazole, dichlorophenyl imidazolodioxalan, which is commercially available from Janssen Pharmaceutica, N.V., under the tradename, "Elubiol", clotrimazole, itraconazole, miconazole, climbazole, tioconazole, sulconazole, butoconazole, fluconazole, miconazole nitrate and any possible stereo isomers and derivatives thereof; piroctone olamine (Octopirox); selenium sulfide; ciclopirox olamine; anti-psoriasis agents such as vitamin D analogs, e.g. calcipotriol, calcitriol, and tacalcitol; vitamin A analogs such as esters of vitamin A, e.g. vitamin A palmitate, retinoids, retinols, and retinoic acid; corticosteroids such as hydrocortisone, clobetasone, butyrate, clobetasol propionate and mixtures thereof.

The amount of benefit agent to be combined with the cleansing composition or the emulsion may vary depending upon, for example, the ability of the benefit agent to penetrate through the skin, hair or nail, the specific benefit agent chosen, the particular benefit desired, the sensitivity of the user to the benefit agent, the health condition, age, and skin, hair, and/or nail condition of the user, and the like. In sum, the benefit agent is used in a "safe and effective amount," which is an amount that is high enough to deliver a desired skin, hair or nail benefit or to modify a certain condition to be treated, but is low enough to avoid serious side effects, at a reasonable risk to benefit ratio within the scope of sound medical judgment. Unless otherwise expressed herein, typically the benefit agent is present in the cleansing system in an amount, based upon the total weight of the system, from about 0.01

percent to about 20 percent, and preferably from about 0.01 percent to about 5.0 percent, and more preferably from about 0.01 percent to about 2.0 percent.

Optionally, commercially available detergent thickeners that are capable of imparting the appropriate viscosity to conditioning shampoo compositions are suitable for use in this invention. If used, the detergent thickeners should be present in the shampoo compositions in an amount sufficient to raise the Brookfield viscosity of the composition to a value of between about 500 to about 10,000 centipoise. Examples of suitable detergent thickeners nonexclusively include: mono or diesters of polyethylene glycol of formula IV.



wherein z is an integer from about 3 to about 200;

fatty acids containing from about 16 to about 22 carbon atoms; fatty acid esters of ethoxylated polyols; ethoxylated derivatives of mono and diesters of fatty acids and glycerine; hydroxyalkyl cellulose; alkyl cellulose; hydroxyalkyl alkyl cellulose; and mixtures thereof. More specifically, suitable detergent thickeners nonexclusively include behenalkonium chloride; cetyl alcohol, quaternium-46, hydroxyethyl cellulose, cocodimonium chloride, polyquaternium-6, polyquaternium-7, quaternium-18, PEG-18 glycerol oleate/cocoate, a mixture of acrylates/steareth-50 acrylate copolymer, laureth-3 and propylene glycol, which is commercially available from Goldschmidt under the tradename "Antil 208," a mixture of cocamidopropylbetaine and glyceryl laurate which is commercially available from Goldschmidt under the tradename, "Antil HS60," a mixture of propylene glycol, PEG 55, and propylene glycol oleate, which is commercially available from Goldschmidt under the tradename, "Antil 414 liquid," and mixtures thereof. Preferred detergent thickeners include polyethylene glycol ester, and more preferably PEG-150 distearate which is available from the Stepan Company of Northfield, Illinois or from Comiel, S.p.A. of Bologna, Italy under the tradename, "PEG 6000 DS".

The above described cleansing composition and cleaning system may be prepared by combining the desired components in a suitable container and mixing them under ambient conditions in any conventional mixing means well known in the art, such as a mechanically stirred propeller, paddle, and the like.

In another preferred embodiment of the cleaning system of the present invention wherein a polymeric emulsifier such as, for example, polyethylene glycol-30 dipolyhydroxystearate (hereinafter "PEG 30") or dimethicone copolyol, are used and water is used as the vehicle, an oil-in-water emulsion may be produced. Although both the PEG 30 and dimethicone copolyol are marketed for use in formulating water-in-oil compositions, we have unexpectedly found that oil-in-water emulsions may be created due to the unique processing steps and conditions employed herein. More specifically, we found that when a

thickening agent, preferably a hydrophilic thickening agent, is neutralized in the hydrophilic phase of the present invention comprising a polymeric emulsifier prior to adding the lipophilic phase of the present invention thereto, the resulting emulsion is in the form of a water-in-oil emulsion. Conversely, when a thickening agent, preferably a hydrophilic thickening agent, is
5 neutralized in the hydrophilic phase of the present invention comprising a polymeric emulsifier after the lipophilic phase of the present invention is added to the hydrophilic phase, the resulting emulsion is unexpectedly in the form of a oil-in-water emulsion.

Cleansing systems of the present invention that are emulsions may contain, based upon the total weight of the emulsion, from about 0.01 percent to about 2 percent, and
10 preferably from about 0.01 percent to about 0.5 percent of hydrophilic thickeners. Suitable neutralizers include any known bases, such as sodium hydroxide, or acids, such as lactic acid, that are capable of neutralizing the hydrophilic thickening agent, in either the hydrophilic phase (if a water-in-oil emulsion is desired) or a mixture of both the hydrophilic phase and the lipophilic phase (if an oil-in-water emulsion is desired) of the present invention to a pH of about
15 5 to about 7 under ambient temperature. In one embodiment, hydrophilic thickeners including acrylates/aminoacrylates copolymer, acrylates/steareth-20 itaconate copolymer, acrylates/ceteth-20 itaconate copolymer, are preferably neutralized with an acid, such as lactic acid. Hydrophilic thickeners including carbomers, modified hydroxyethylcellulose, polyvinylacetate/maleic anhydride (PVA/MA) decadiene crosspolymer, and
20 acrylates/steareth-20 methacrylate copolymer, are preferably neutralized with a base, such as sodium hydroxide (20%).

In one embodiment, the hydrophilic phase may be comprised of one or more of the following components: water, thickener, cleansing enhancer, nonfoaming surfactant, and water dispersible component, and the lipophilic phase may be comprised of one or more of
25 the following components: silicone, ester, and polymeric emulsifier.

We have also surprisingly found that the cleansing system of the present invention possesses good aesthetic properties without causing any significant ocular discomfort to the user. It is well-known in the art that most emulsifiers having a relatively low molecular weight are irritating regardless of their hydrophilic lipophilic balance ("HLB") value. However, we have
30 surprisingly found that when the cleansing system of the present invention is produced using the particular polymeric emulsifiers and/ or thickeners set forth herein, the resulting cleanser is gentle and possesses a low degree of ocular and skin irritation.

Another embodiment of this invention is directed to a foaming composition comprising, based upon the foaming composition, from about 0.1 percent to about 30 percent, e.g. from
35 about 0.1 percent to about 5 percent of a water dispersible component; from about 0.1 percent to about 30 percent, e.g. from about 0.1 percent to about 5 percent of an ester; from

about 1.0 percent to about 98 percent, e.g. from about 30 percent to about 98 percent or from about 45 percent to about 90 percent of water; and from about 2.0 percent to about 20 percent, e.g. from about 5.0 percent to about 15 percent of a foaming surfactant.

Optionally, the foaming composition may also be comprised of one or more of the following components, based upon the total weight of the foaming composition: a) from about 0.1 percent to 5 percent, e.g. from about 0.5 percent to about 1.5 percent of a polymeric emulsifier, a thickener, or mixture thereof; b) from about 0.1 percent to about 5 percent, e.g. from about 1 percent to about 3 percent of a cleaning enhancer; c) from about 0.001 percent to about 20 percent of a benefit agent; and d) from about 0.1 percent to about 30 percent, e.g. from about 0.1 percent to about 5 percent of a liquid silicone.

Another embodiment of the present invention is directed to a method for depositing a benefit agent onto the skin, hair and/or nails comprised of applying either the above-described cleansing system or cleansing composition with an effective amount of a benefit agent to a desired location on a human or animal. While the frequency and amount of the benefit agent-containing cleaning system to be applied will depend upon, for example, the type and amount of benefit agent available, the intended usage of the final composition, i.e. therapeutic versus maintenance regimen, the amount and type of detergent present, and the sensitivity of the individual user to the composition/emulsion, typically the benefit agent-containing cleaning system of the present invention should be topically applied to affected body parts at regular intervals, and preferably from about 2 to about 14 times per week. More preferably, the composition/emulsion is applied more frequently during the initial stages of treatment, e.g. from about 5 to about 7 times per week until the desired effect is achieved, then less frequently when maintenance is desired, e.g. from about 2 to about 5 times per week.

We have unexpectedly found that the above-described cleansing composition and cleansing system are capable of efficiently mediating the deposition and permeation of various benefit agents, such as antidandruff agents, onto and into the skin following topical administration thereto. More specifically, we have surprisingly found that when benefit agents are combined with either the cleansing composition or the cleaning system of the present invention, the amount of benefit agents deposited onto and/or into the skin, hair, and/or nails is about 50% greater than the amount of benefit agents deposited onto and/or into the skin, hair, and/or nails after application of known, commercial benefit agent-containing cleansers.

An alternative preferred embodiment of the present invention is directed to a method for treating hair loss, such as hair loss resulting from alopecia, comprising topically applying the above-described cleaning system and the hair loss benefit agent to a desired location on an animal or human, wherein the benefit agent is comprised of an effective amount of a hair

loss treatment agent such as minoxidil or mixture thereof. As used herein, "hair loss treatment agents" shall include agents capable of growing hair and/or agents capable of preventing the loss of hair. By "effective amount," it is meant an amount effective for treating hair loss and preferably may range from, based upon the total weight of the cleansing system, from about 0.001 percent to about 20 percent, and preferably from about 1 percent to about 5 percent.

Examples of benefit agents suitable for treating hair loss include, but are not limited to potassium channel openers or peripheral vasodilators such as minoxidil, diazoxide, and compounds such as N'-cyano-N-(tert-pentyl)-N'-3-pyridinyl-guanidine ("P-1075") as disclosed in United States Patent No.: 5,244,664, which is incorporated herein by reference; vitamins, such as vitamin E and vitamin C, and derivatives thereof such as vitamin E acetate and vitamin C palmitate; hormones, such as erythropoietin, prostaglandins, such as prostaglandin E₁ and prostaglandin F₂-alpha; fatty acids, such as oleic acid; diuretics such as spironolactone; heat shock proteins ("HSP"), such as HSP 27 and HSP 72; calcium channel blockers, such as verapamil HCL, nifedipine, and diltiazemamiloride; immunosuppressant drugs, such as cyclosporin and Fk-506; 5 alpha-reductase inhibitors such as finasteride; growth factors such as, EGF, IGF and FGF; transforming growth factor beta; tumor necrosis factor; non-steroidal anti-inflammatory agents such as benoxaprofen; retinoids such as tretinoin; cytokines, such as IL-6, IL-1 alpha, and IL-1 beta; cell adhesion molecules such as ICAM; glucocorticoids such as betametasone; botanical extracts such as aloe, clove, ginseng, rehmannia, swertia, sweet orange, zanthoxylum, *Serenoa repens* (saw palmetto), *Hypoxis rooperi*, stinging nettle, pumpkin seeds, and rye pollen; other botanical extracts including sandalwood, red beet root, chrysanthemum, rosemary, burdock root and other hair growth promoter activators which are disclosed in DE 4330597 which is incorporated by reference in its entirety herein; homeopathic agents such as *Kalium Phosphoricum D2*, *Azadirachta indica D2*, and *Joborandi DI*; genes for cytokines, growth factors, and male-patterned baldness; antifungals such as ketoconazole and elubiol; antibiotics such as streptomycin; proteins inhibitors such as cycloheximide; acetazolamide; benoxaprofen; cortisone; diltiazem; hexachlorobenzene; hydantoin; nifedipine; penicillamine; phenothiazines; pinacidil; psoralens, verapamil; zidovudine; alpha-glucosylated rutin having at least one of the following rutins: quercetin, isoquercitrin, hespeddin, naringin, and methylhesperidin, and flavonoids and transglycosidated derivatives thereof which are all disclosed in JP 7002677, which is incorporated by reference in its entirety herein; and mixtures thereof.

Preferred hair loss treatment agents include minoxidil, 6-(1-piperidinyl)-2,4-pyrimidinediamine-3-oxide, N'-cyano-N-(tert-pentyl)-N'-3-pyridinyl-guanidine, finasteride, retinoids and derivatives thereof, ketoconazole, elubiol or mixtures thereof.

Another embodiment of the present invention is directed to a method for inhibiting hair growth comprising topically applying the above-described composition/system combined with a benefit agent to a desired area on an animal or human for inhibiting hair growth, wherein the benefit agent is comprised of an effective amount of a hair growth inhibiting agent. In a preferred embodiment, the cleaning system contains, based upon the total weight of the cleaning system, from about 0.001 percent to about 20 percent, and preferably from about 0.01 percent to about 5 percent hair growth inhibiting agent.

Examples of benefit agents suitable for use in inhibiting hair growth include: serine proteases such as trypsin; vitamins such as alpha-tocophenol (vitamin E) and derivatives thereof such as tocophenol acetate and tocophenol palmitate; antineoplastic agents, such as doxorubicin, cyclophosphamide, chlormethine, methotrexate, fluorouracil, vincristine, daunorubicin, bleomycin and hydroxycarbamide; anticoagulants, such as heparin, heparinoids, coumaerins, detran and indandiones; antithyroid drugs, such as iodine, thiouracils and carbimazole; lithium and lithium carbonate; interferons, such as interferon alpha, interferon alpha-2a and interferon alpha-2b; retinoids, such as retinol (vitamin A), isotretinoin; glucocorticoids such as betamethasone, and dexamethosone; antihyperlipidaemic drugs, such as triparanol and clofibrate; thallium; mercury; albendazole; allopurinol; amiodarone; amphetamines; androgens; bromocriptine; butyrophenones; carbamazepine; cholestyramine; cimetidine; clofibrate; danazol; desipramine; dixyrazine; ethambutol; etionamide; fluoxetine; gentamicin, gold salts; hydantoins; ibuprofen; imipramine; immunoglobulins; indandiones; indomethacin; intraconazole; levodopa; maprotiline; methysergide; metoprolol; metyrapone; nadolol; nicotinic acid; potassium thiocyanate; propranolol; pyridostimine; salicylates; sulfasalazine; terfenadine; thiamphenicol; thiouracils; trimethadione; tropanolol; valproic acid; and mixtures thereof.

Preferred hair growth inhibitory agents include serine proteases, retinol, isotretinoin, betamethoisone, alpha-tocophenol and derivatives thereof, or mixtures thereof.

Another preferred embodiment of the present invention is directed to a method for treating acne and for reducing the signs of aging, i.e. wrinkles, fine lines, and other manifestations of photodamage, comprising topically applying the above-described cleaning system and the relevant benefit agent to the skin of an animal or human at a desired area, wherein the benefit agent is comprised of an effective amount of an anti-acne agent or an anti-aging agent, respectively.

Examples of suitable anti-aging agents include, but are not limited to inorganic sunscreens such as titanium dioxide and zinc oxide; organic sunscreens such as octyl-methoxy cinnamates and derivatives thereof; retinoids; vitamins such as vitamin E, vitamin A, vitamin C, vitamin B, and derivatives thereof such as vitamin E acetate, vitamin C palmitate, and the like; antioxidants including beta carotene, alpha hydroxy acids such as

glycolic acid, citric acid, lactic acid, malic acid, mandelic acid, ascorbic acid, alpha-hydroxybutyric acid, alpha-hydroxyisobutyric acid, alpha-hydroxyisocaproic acid, atrolactic acid, alpha-hydroxyisovaleric acid, ethyl pyruvate, galacturonic acid, glucoheptonic acid, glucoheptono 1,4-lactone, gluconic acid, gluconolactone, glucuronic acid, glucuronolactone, glycolic acid, isopropyl pyruvate, methyl pyruvate, mucic acid, pyruvic acid, saccharic acid, saccaric acid 1,4-lactone, tartaric acid, and tartronic acid; beta hydroxy acids such as beta-hydroxybutyric acid, beta-phenyl-lactic acid, beta-phenylpyruvic acid; botanical extracts such as green tea, soy, milk thistle, algae, aloe, angelica, bitter orange, coffee, goldthread, grapefruit, hoellen, honeysuckle, Job's tears, lithospermum, mulberry, peony, pueraria, nice, safflower, and mixtures thereof.

Preferred anti-aging agents include retinoids, anti-oxidants, alpha-hydroxy acids and beta-hydroxy acid with retinol and tretinoin being most preferred.

Suitable amounts of anti-aging agents include, based upon the total weight of the described cleaning system, from about 0.01 percent to about 20 percent, and preferably from about 0.04 percent to about 5 percent.

Examples of suitable anti-acne agents include, but are not limited to topical retinoids (tretinoin, isotretinoin, motretinide, adapalene, tazarotene, azelaic acid, retinol); salicylic acid; benzoyl peroxide; resorcinol; antibiotics such as tetracycline and isomers thereof, erythromycin, and the anti-inflammatory agents such as ibuprofen, naproxen, hetprofen; botanical extracts such as alnus, arnica, artemisia capillaris, asiasarum root, birrh, calendula, chamomile, cnidium, comfrey, fennel, galla rhois, hawthorn, houttuynia, hypericum, jujube, kiwi, licorice, magnolia, olive, peppermint, philodendron, salvia, sasa albo-marginata; imidazoles such as ketoconazole and elubiol, and those described in Gollnick, H et al. 196(I) Dermatology Sebaceous Glands, Acne and Related Disorders, 119-157 (1998), which is incorporated by reference herein, and mixtures thereof.

Preferred anti-acne agents include benzoyl peroxide, retinol, elubiol, antibiotics, and salicylic acid, with retinol and tretinoin being most preferred.

Suitable amount of anti-acne agents include, based upon the total weight of the described cleaning system, from about 0.01 percent to about 10 percent, and preferably from about 0.04 percent to about 5 percent.

Another preferred embodiment of the present invention is directed to a method for depigmenting the skin, comprising topically applying to skin at a desired area the above-described cleaning system and an effective amount of the depigmentation benefit agent. Suitable effective amounts of depigmentation agents include, based upon the total weight of the described cleaning system, from about 0.01 percent to about 10 percent, and preferably from about 0.04 percent to about 5 percent.

Examples of suitable depigmentation agents include, but are not limited to soy and derivatives thereof, retinoids such as retinol; Kojic acid and its derivatives such as, for example, kojic dipalmitate; hydroquinone and its derivatives such as arbutin; tranexamic acid; vitamins such as niacin, vitamin C and its derivatives; azelaic acid; placertia; licorice; 5 extracts such as chamomile and green tea, and mixtures thereof, with retinol, Kojic acid, and hydroquinone, being preferred.

An alternative preferred embodiment of the present invention is directed to a method for treating the symptoms and/or the diseases of dandruff, seborrheic dermatitis and/or psoriasis, comprising topically applying the above-described cleaning system and the 10 relevant benefit agent to a location desired wherein the benefit agent is comprised of an effective amount of a dandruff treatment agent, a seborrheic dermatitis treatment agent, or a psoriasis treatment agent, respectively. As used herein, "dandruff treatment agent," "seborrheic dermatitis treatment agent," or a "psoriasis treatment agent," respectively, shall include agents capable of treating the symptoms and/or the diseases of dandruff, seborrheic 15 dermatitis, and psoriasis, respectively. By "effective amount," it is meant an amount effective for treating the disease and/or the symptoms associated therewith and preferably may range from, based upon the total weight of the cleaning system, from about 0.001 percent to about 10 percent, and preferably from about 0.01 percent to about 5 percent.

Examples of benefit agents suitable for treating the symptoms and/or the diseases 20 of dandruff, seborrheic dermatitis and/or psoriasis, respectively, nonexclusively include those set forth above with shale oil and derivatives thereof, elubiol, ketoconazole, coal tar, salicylic acid, zinc pyrithione, selenium sulfide, hydrocortisone, sulfur, menthol, pramoxine hydrochloride, and mixtures thereof being particularly preferred.

The compositions of the present invention may be directed applied to the skin or may 25 be applied onto other delivery implements such as wipes, sponges, brushes, and the like. The compositions may be used in products designed to be left on the skin, wiped from the skin, or rinsed off of the skin.

The invention illustratively disclosed herein suitably may be practiced in the absence of any component, ingredient, or step which is not specifically disclosed herein. Several 30 examples are set forth below to further illustrate the nature of the invention and the manner of carrying it out. However, the invention should not be considered as being limited to the details thereof.

Examples

Example 1: Preparation of Oil in Water Emulsion Incorporating a Polymeric Emulsifier**Preparation of Lipophilic Phase:**

11 g of PEG-30 dipolyhydroxystearate, available from Uniqema, Inc. under the tradename "Arlacel P-135," 50 g of isononyl isononanoate, available from Alzo, Inc. under the tradename, "Wickenol 151," 50 g of a mixture of hexyldecyl benzoate and butyloctyl benzoate, available from C.P. Hall Company under the tradename, "Hallstar AB," and 50 g of cyclomethicone available from Dow Corning under the tradename, "Dow 344 Fluid" were combined into a glass beaker containing a propeller stirrer and heated to a temperature of 60 °C until homogeneous.

Preparation of Hydrophilic Phase:

Into a primary glass beaker containing 795 g of deionized water, 5 g of PEG-8 caprylic/capric glycerides available from Trivent Inc. under the tradename, "Trivasol BW" was added thereto with stirring at 25°C until homogeneous. For aiding in dispersion of the thickener in the formulation, 4 g of carbomer available from B.F. Goodrich, Inc. under the tradename, "Carbopol Ultrez" were added to 30 g of dimethylisosorbide available from Uniqema, Inc. under the tradename, "Arlasolve DMI" in a separate beaker with hand stirring. Into the dimethylisosorbide mixture was then added 2 g. of methylparaben and 1 g of propylparaben with hand stirring until homogeneous to produce a pre-mixture. The pre-mixture was then added to the primary glass beaker with constant stirring until the resulting mixture was homogeneous.

Preparation of Final Composition:

The lipophilic phase was then added to the hydrophilic phase with constant stirring at 25°C until homogeneous. 2 g of triethanolamine available from Union Carbide under the tradename, "Trolamine 99%" was then added to the resulting mixture with stirring until homogeneous.

Example 2: Preparation of Polymeric Emulsifier-Free Oil in Water Emulsion**Preparation of Lipophilic Phase:**

20 g of isostearyl palmitate, available from Brooks Industries, under the tradename "Loronate OP," 20 g of isononyl isononanoate, available from Alzo, Inc. under the tradename, "Wickenol 151," 20 g of cetyl octanoate, available from Brooks Industries, under the tradename "Loronate CIO," 20 g of pentaerythritol tetraoctanoate available from Brooks Industries, under the tradename "Loronate PT," and 20 g of cyclomethicone available from Dow Corning under the tradename, "Dow 345 Fluid" were combined into a glass beaker at a temperature of 25 °C and stirred until homogeneous.

Preparation of Hydrophilic Phase:

Into a primary glass beaker containing 859.7 g of deionized water, 5 g of carbomer available from B.F. Goodrich, Inc. under the tradename, "Carbopol Ultrez" was added thereto with stirring at a temperature of 25 °C until homogenous. Into a separate beaker was added 7.5 g. of sucrose cocoate available from Croda, Inc. under the tradename, "Crodesta SL-40," 7.5 g. of PEG –6 Capric/caprylic glycerides available from Croda, Inc. under the tradename, "Glycerox 767," 10 g of hexylene glycol, 3 g. of methylparaben and 0.5 g of propylparaben with hand stirring until homogeneous to produce a pre-mixture. The pre-mixture was then added to the primary glass beaker with constant stirring until the resulting mixture was homogeneous.

Preparation of Final Composition:

After the lipophilic phase was added to the hydrophilic phase with constant stirring at 25°C until homogeneous, 6.8 g. of a 20% aqueous solution of sodium hydroxide was then added thereto with stirring at a temperature of 25 °C. The resulting mixture was then mixed for 15 minutes.

Example 3: Use of Emulsions to Remove Make-Up

Onto 3 respective ½ inch by 1 inch rectangle areas located on the forearms of caucasian women were applied each of the make-ups described in Table A below:

Table A: Types of Make-Up

<u>Type of Make-up</u>	<u>Supplier</u>	<u>Tradename / Color</u>
Foundation	Revlon	"Colorstay Make-Up" – in "Mocha"

Lipstick	Estee Lauder	"All Day Lipstick" – in "Regal Red"
Mascara	Revlon	"Colorstay Lashcolor Waterproof" - in "Navy"

After dipping a wooden applicator from Scientific Products into the foundation, the foundation was spread throughout the designated area and evenly distributed therein with a finger cot.

- 5 The lipstick was directly applied to the designated area and evenly distributed therein with a finger cot.

The mascara was spread with its wand in the designated rectangular area and evenly distributed therein with a finger cot.

- 10 After the make-up dried for 30 minutes at room temperature, 0.8 ml of the emulsion of Example 1 was dispensed via syringe across all three test sites. Using a wooden tongue depressor, the emulsion was rubbed across all three test sites in a rapid, circular motion for 30 seconds. Then, each of the three test sites were gently wiped with a Kim-wipe.

- This procedure was repeated but wherein the emulsion was replaced with the following respective cleansing products: a) "Galenic," which is available from Laboratoires Galenic, a
15 division of PIERRE FABRE, under the tradename, "Lait Demaquillant Hydratant (Moisturizing Cleansing Milk);" b) "Cetaphil," which is available from GALDERMA INTERNATIONAL under the tradename, "Lotion Nettoyante/Haute Tolerance (pour peaux sensibles);" c) "pH 5.5 3-in-1," which is available from JOHNSON & JOHNSON LIMITED under the tradename, "JOHNSON's pH 5.5 – 3 in 1 Facial Cleanser;" d) "Vichy," which is available from VICHY
20 LABORATORIES under the tradename, "Demaquillant Integral (a l'eau thermale apaisante);" e) "Oil of Olay," which is commercially available from THE PROCTER & GAMBLE COMPANY under the tradename, "Oil of Olay Facial Cleansing Lotion;" and f) "F # 8626-015," which is similar to the formulation as set forth in Example 2, except as follows in Table B below:

Table B: Contrast of Example 2 formulation with formula 8626-015:

Example 2 Formulation	Modified Example 2 (F # 8626-015)	Component
0.3% (3 grams)	0.2% (2 grams)	methylparaben
0.05% (0.5 grams)	0.1% (1 gram)	Propylparaben

85.97% (859.7 grams)	86.02% (860.2 grams)	Water
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Photographs of the test sites were obtained after the make-up was applied and after the three sites were rubbed with the above six products as shown in FIGURE 1.

As evidenced in FIGURE 1, the emulsion of modified Example 2 was superior with respect to removal of mascara, foundation, and lipstick in comparison to the other cleansers. The emulsion of Example 1 (not shown in FIGURE 1) also was effective in removing all three types of make-up. This Example showed that the emulsions of the present invention are superior with respect to known commercial cleansers in removing a variety of make-up types from the skin.

Example 4: Preparation of Shower Gel Containing Sunscreen

A composition containing the following ingredients as set forth in Table C is prepared as follows:

Table C: Composition of Shower Gel Containing Sunscreen

<u>Tradename</u>	<u>Chemical Name</u>	<u>Weight (g)</u>	<u>Supplier</u>
WATER PHASE			
-----	Deionized water	696.5	-----
-----	Methylparaben	3	-----
Comperian 100K	Cocamide MEA	30	Henkel Corporation
Planteren 1200N	Lauryl glucoside	50	Henkel Corporation
Schercomid HT 60	PEG-50-Tallow Amide	50	Scher Chemical Corporation
	Hexylene glycol	10	
Standamox CAW	Cocamidopropylamine oxide	50	Henkel Corporation
OIL PHASE			
Hallstar AB	Hexyldecyl benzoate & butyloctyl benzoate	50	C.P. Hall Company
Ariacel P-135	PEG-30 Dipolyhydroxystearate	30	Uniqema, Inc.
Neo Heliopan	Menthyl anthranilate	30	Haarmann & Reimer
POST ADDITION			
Crothix Liquid	PEG 150 pentaerythrityl tetrastearate & PEG-6 caprylic/capric glycerides & water	10	Croda Inc.
Kathon CG	Methylchloroisothiazolinone & methylisothiazolinone	0.5	Rohm & Haas

After heating the deionized water to a temperature of about 75 °C to about 80 °C in a beaker, the hexylene glycol followed by the methylparaben is added thereto with mixing until solubilized. The cocamide MEA followed by the lauryl glucoside is then added thereto sequentially at constant temperature. The PEG-50 tallow amide is then melted and added to the resulting mixture at constant temperature. The cocamidopropylamine oxide is then added thereto at constant temperature to form a water phase premixture.

In another beaker, all of the oil phase components are combined with mixing at a temperature of about 75 °C to about 80 °C until uniform to form an oil phase premixture. The oil phase premixture is then added to the water phase premixture at constant temperature.

10 The "Crothix Liquid" component is then added thereto with mixing until uniform, then the temperature is lowered to about 50 °C. The "Kathon CG" component is then added thereto, and the resulting mixture is cooled to room temperature.

The resulting cleansing formulation will possess excellent foaming properties and, after rinsing, will leave the skin with a light, moisturized feeling. Moreover, an effective amount of sunscreen agent will be deposited on the skin after rinsing the formulation from the skin with water.

Examples 5 and 6: Preparation of Foaming Anti-Acne Shower Gels

The composition of Example 5 containing the following ingredients as set forth in Table D is prepared as follows:

Table D: Compositions of Foaming Shower Gel

Tradename	Chemical Name	Wt % Ex. 5	Wt % Ex. 6	Supplier
WATER PHASE				
-----	Deionized water	49.440	49.654	-----
-----	Methylparaben	0.3	0.3	-----
Comperian 100K	Cocamide MEA	3.0	3.0	Henkel Corporation
Planteren 1200N	Lauryl glucoside	5.0	5.0	Henkel Corporation
Schercomid HT 50	PEG-50 Tallow Amide	5.0	5.0	Scher Chemical Co.
Standamox CAW	Cocamidopropylamine oxide	5.0	5.0	Henkel Corporation
Schercoquat DAS	Quaternium 61	2.0	2.0	Scher Chemical Co.
Jaguar C17	Guar hydroxypropyl trimonium chloride	0.5	0.5	Rhodia

	Glycerin	5.0	5.0	----
	Deionized water	10	10	----
	Citric acid (20% soln)	0.566	0.490	----
OIL PHASE				
Abil EM 90	Cetyl dimethicone copolyol	0.0	3.0	Goldschmidt Chemical Corporation
Arlacel P-135	PEG-30 Dipolyhydroxystearate	3.0	0.0	Unichema, Inc.
Wickenol 151	Isononyl isononanoate	1.0	1.0	Alzo
Volpo L3 Special	Laureth – 3	5.0	5.0	Croda, Inc.
POST ADDITION				
Arlasolve DMI	Dimethyl isosorbide	4.0	4.0	Croda Inc.
Salicylic acid	Salicylic acid	2.0	2.0	----
NaOH (10%)	Sodium hydroxide	0.144	none	----
Kathon CG	Methyl-chloroisothiazolinone & methylisothiazolinone	0.050	0.050	Rohm & Haas

Preparation of Hydrophilic Phase

The water phase premixture is prepared in accordance with Example 3, but with the following additional steps: The quaternium-61 is added with mixing to the mixture after the
5 addition of the cocamidopropyl amine oxide thereto at constant temperature until homogeneous.

In a separate glass beaker equipped with a propeller, the guar hydroxypropyl trimonium chloride and the glycerin are combined with stirring. The additional deionized water is then slowly added thereto with constant stirring. The resulting mixture is then mixed for
10 about 15 minutes, then acidified with the citric acid solution. The acidified solution is then added to the water phase premixture with stirring at a temperature of about 75 °C to about 80 °C until homogeneous.

Preparation of Oil Phase

PEG-30 dipolyhydroxystearate is combined with the laureth-3 in another beaker with
15 mixing at a temperature of about 75 °C to about 80 °C to form an oil phase premixture.

The oil phase premixture is then added to the water phase premixture with mixing at constant temperature.

Preparation of Final Composition:

The lipophilic phase is added to the hydrophilic phase with constant stirring at 75°C
20 until homogenous. After the temperature of the resulting mixture is lowered to 50°C with

continuous stirring, a premixed solution of 40 grams of Dimethyl Isosorbide available from Uniqema Inc. under the tradename "Arlasolve DMI" and 20 grams of salicylic acid are added thereto with stirring. After the temperature of the resulting mixture is lowered to 30°C, its pH is adjusted to approximately 3.1 by adding 1.4 grams of a 10% aqueous sodium hydroxide solution with continued stirring until the temperature of the resulting mixture reaches 25°C.

This process is repeated for the formulation of Example 6, but the PEG-30 dipolyhydroxystearate is replaced with cetyl dimethicone copolyol.

The above formulations will result in gels that possess good foaming properties as well as leave the skin with a pleasant, lightly moisturized "after feel" due to the effective deposition of the humectant. Similarly, the formulations will be useful as anti-acne formulations due to their ability to effectively deposit the salicylic acid component into and onto the skin.

Example 7: Preparation of Foaming Gel Containing Sunscreen

Preparation of Hydrophilic Phase:

Into a primary glass beaker equipped with a propeller containing 667 grams of deionized water, 30 grams of Hexylene Glycol available from Shell Chemical Company and 3 grams of methylparaben are added thereto stirring at 75°C until homogenous. While maintaining the agitation and temperature as constant, 30 grams of cocomonoethanolamide (100% active) available from Henkel Corporation under the tradename "Comperian 100K", 50 grams of Lauryl Glucoside (50% active) available from Henkel Corporation under the tradename "Plantaren 1200N", 50 grams of PEG-50 Tallow Amide (100% active) available from Scher Chemical Corporation under the tradename "Schercomid HT 60", and 50 grams of Cocamidopropylamine Oxide (30% active) available from Henkel Corporation under the tradename "Standamox CAW" are added to this aqueous mixture and mixed until homogenous.

Preparation of Lipophilic Phase:

50 grams of Hexyldecyl Benzoate and Butyloctyl Benzoate available from C.P. Hall Co. under the tradename "HallStar AB", 30 grams of PEG-30 Dipolyhydroxystearate available from Unichema, Inc. under the tradename "Arlacel P135" and 30 grams of Menthyl Anthranilate available from Haarmann & Reimer Corp. under the tradename "Neo Heliopan MA" are combined into another glass beaker equipped with a propeller and stirred until homogenous with heat until the resulting lipophilic mixture was at a temperature of 75°C.

Preparation of Final Composition:

The lipophilic phase is then added to the hydrophilic phase with constant stirring at 75°C. 10 grams of a mixture of PEG-150 Pentaerythrityl Tetrastearate, PEG-6 Caprylic/Capric Glycerides, and Water available from Croda Inc. under the tradename "Crothix Liquid" is then added thereto. The resulting mixture is then stirred at 75°C until homogenous, then cooled to room temperature with constant stirring.

The resulting cleansing formulation possesses excellent foaming properties and, after rinsing, leaves the skin with a light, moisturized feeling. The formulation also effectively deposits the sunscreen agent into and onto the skin.

10

Example 8: Preparation of Foaming Acne WashPreparation of Hydrophilic Phase:

Into a primary glass beaker equipped with a propeller stirrer containing 414.9 grams of deionized water, 3 grams of methylparaben are added thereto at 75°C until homogenous. While maintaining constant agitation and temperature, 30 grams of cocomonoethanolamide (100% active) available from Henkel Corporation under the tradename "Comperlan 100K", 50 grams of Lauryl Glucoside (50% active) available from Henkel Corporation under the tradename "Plantaren 1200N", 50 grams of PEG-50 Tallow Amide (100% active) available from Scher Chemical Corporation under the tradename "Schercomid HT 60", 50 grams of Cocamidopropylamine Oxide (30% active) available from Henkel Corporation under the tradename "Standamox CAW", and 20 grams of Quaternium-61 (90% active) available from Scher Chemical Corporation under the tradename "Schercoquat DAS" are added to this aqueous mixture and mixed until homogenous to form a primary hydrophilic phase.

5 grams of Guar Hydroxypropyl Trimonium Chloride available from Rhodia Inc. under the tradename "Jaguar C17" is combined with stirring with 50 grams of glycerin into a separate glass beaker equipped with a propeller stirrer. 100 grams of deionized water is slowly added with mixing to this mixture at constant temperature. After the resulting mixture is mixed for 15 minutes until homogenous, the mixture is acidified with 5.7 grams of a 20% aqueous solution of citric acid. After the acidified mixture is added to the primary hydrophilic phase, the resulting hydrophilic mixture is stirred at 75°C until homogenous.

Preparation of Lipophilic Phase:

50 grams of Isopropyl PPG-2-Isodeceth-7 Carboxylate available from Clariant Corporation under the tradename "Velsan D8P-3", 30 grams of PEG-30 Dipolyhydroxystearate available from Unichema, Inc. under the tradename "Arlacel P135" and 30 grams of Laureth-3 (100% active) available from Croda Inc. are combined with

stirring into a glass beaker equipped with a propeller stirrer until homogenous and heated to a temperature of 75°C.

Preparation of Final Composition:

- 5 The lipophilic phase is added to the hydrophilic phase with constant stirring at 75°C until homogenous. After the temperature of the resulting mixture is lowered to 50°C with continuous stirring, a premixed solution of 40 grams of Dimethyl Isosorbide available from Unichema Inc. under the tradename "Arlasolve DMI" and 20 grams of salicylic acid is added thereto with stirring. After the temperature of the resulting mixture is lowered to 30°C, its pH
10 is adjusted to approximately 3.1 by adding 1.4 grams of a 10% aqueous sodium hydroxide solution with continued stirring until the temperature of the resulting mixture reaches 25°C.

The resulting wash product will not only possess excellent foaming properties but will also be an effective anti-acne product due to its ability to deposit the anti-active agent into and onto the skin.

15

Example 9: Preparation of Water-in-Oil Emulsion

Preparation of Lipophilic Phase:

- 20 g of isostearyl palmitate, available from Brooks Industries, under the tradename
20 "Loronate OP," 20 g of isononyl isononanoate, available from Alzo, Inc. under the tradename, "Wickenol 151," 20 g of cetyl octanoate, available from Brooks Industries, under the tradename "Loronate CIO," 20 g of pentaerythritol tetraoctanoate available from Brooks Industries, under the tradename "Loronate PT," and 20 g of cyclomethicone available from Dow Corning under the tradename, "Dow 345 Fluid" were combined into a glass beaker at a temperature of 25 °C
25 and stirred until homogeneous.

Preparation of Hydrophilic Phase:

- Into a primary glass beaker containing 859.7 g of deionized water, 5 g of carbomer available from B.F. Goodrich, Inc. under the tradename, "Carbopol Ultrez" was added thereto with stirring at a temperature of 25 °C until homogenous. Into a separate beaker was added
30 7.5 g. of sucrose cocoate available from Croda, Inc. under the tradename, "Crodesta SL-40," 7.5 g. of PEG -6 Capric/caprylic glycerides available from Croda, Inc. under the tradename,

"Glycerox 767," 10 g of hexylene glycol, 3 g. of methylparaben and 0.5 g of propylparaben with hand stirring until homogeneous to produce a pre-mixture. The pre-mixture was then added to the primary glass beaker with constant stirring until the resulting mixture was homogeneous.

Preparation of Final Composition:

- 5 After the 6.8 g. of a 20% aqueous solution of sodium hydroxide was added to the hydrophilic phase with constant stirring at 25°C until homogeneous, the lipophilic phase was added thereto with stirring at a temperature of 25 °C. The resulting mixture was then mixed for 15 minutes.

Example 10: Preparation of Oil-in-Water Emulsion Containing Retinol

10 Preparation of Lipophilic Phase:

- 11 g of PEG-30 dipolyhydroxystearate, available from Uniqema, Inc. under the tradename "Arlacel P-135," 50 g of isononyl isononanoate, available from Alzo, Inc. under the tradename, "Wickenol 151," and 50 g of a mixture of hexyldecyl benzoate and butyloctyl benzoate, available from C.P. Hall Company under the tradename, "Hallstar AB" were
15 combined with continuous mixing in a vessel and heated to a temperature of 45°C until homogeneous. After the mixture was cooled to a temperature of 25 °C, 50 g of cyclomethicone available from Dow Corning under the tradename, "Dow 344 Fluid" and 6.9 g of a mixture of vitamin A alcohol and polysorbate 20 in a 1:1 weight ratio were added thereto with continuous mixing under an Argon blanket and under yellow light into a glass beaker containing a propeller
20 stirrer until homogeneous. All subsequent procedures with this lipophilic phase were conducted under these conditions of argon blanket and yellow light until the formulation is placed into an oxygen and light impermeable container.

Preparation of Hydrophilic Phase:

- Into a primary glass beaker containing 795 g of deionized water, nitrogen was bubbled
25 therein until the subsequent addition of the lipophilic phase thereto so as to minimize exposure to oxygen. 5 g of PEG-8 caprylic/capric glycerides available from Trivent Inc. under the tradename, "Trivasol BW" was then added thereto with stirring at 25°C until homogeneous. For aiding in dispersion of the thickener in the formulation, 4 g of carbomer available from B.F.

Goodrich, Inc. under the tradename, "Carbopol Ultrez" were added to 30 g of dimethylisosorbide available from Uniqema, Inc. under the tradename, "Arlasolve DMI" in a separate beaker with hand stirring. Into the dimethylisosorbide mixture was then added 2 g. of methylparaben and 1 g of propylparaben with hand stirring until homogeneous to produce a pre-mixture. The pre-mixture was then added to the primary glass beaker with constant stirring until the resulting mixture was homogeneous.

Preparation of Final Composition:

The lipophilic phase was then added to the hydrophilic phase with constant stirring at 25°C until homogeneous. 2 g of triethanolamine available from Union Carbide under the tradename, "Trolamine 99%" was then added to the resulting mixture with stirring until homogeneous. The final emulsion contains the components as set forth in Table E:

Table E: Emulsion Components

Chemical Name	Trade Name	%(wt/wt)
PEG-30 dipolyhydroxystearate	Arlacel P-135	1.1
Isononyl isononanoate	Wickenol	5.0
Hexyldecyl benzoate and butyloctyl benzoate	Hallstar AB	5.0
Cyclomethicone	Dow 344 Fluid	5.0
Vitamin A alcohol and Tween 20	Retinol 50C	0.69
Water	Water	78.81
Carbomer	Carbopol Ultrez	0.40
PEG-8 caprylic/capric glycerides	Trivasol BW	0.50
Methylparaben	Methylparaben	0.20
Propylparaben	Propylparaben	0.10
Dimethyl isosorbide	Arlasolve DMI	3.0
triethanolamine	Trolamine 99%	0.2

Example 11: Preparation of Water-in-Oil Emulsion Containing Retinol

Preparation of Lipophilic Phase:

11 g of PEG-30 dipolyhydroxystearate, available from Uniqema, Inc. under the tradename "Arlacel P-135," 30 g of isononyl isononanoate, available from Alzo, Inc. under the tradename, "Wickenol 151," and 30 g of a mixture of hexyldecyl benzoate and butyloctyl benzoate, available from C.P. Hall Company under the tradename, "Hallstar AB" were combined in a vessel with mixing and heated to a temperature of 45 °C until homogeneous. After the resulting mixture was cooled to a temperature of 25°C, 30 g of cyclomethicone available from Dow Corning under the tradename, "Dow 344 Fluid" and 6.9 g of a mixture of vitamin A alcohol and polysorbate 20 in a 1:1 weight ratio were added thereto with continuous mixing under an Argon blanket and under yellow light into a glass beaker containing a propeller stirrer until homogeneous. All subsequent procedures with this lipophilic phase was conducted under these conditions of argon blanket and yellow light until the formulation is placed into an oxygen and light impermeable container.

Preparation of Hydrophilic Phase:

Into a primary glass beaker containing 863.2 g of deionized water, nitrogen was bubbled therein in order to eliminate dissolved oxygen contained therein. The nitrogen continued to be bubbled therein until the subsequent addition of the lipophilic phase thereto. 5 g of PEG-8 caprylic/capric glycerides available from Trivent Inc. under the tradename, "Trivasol BW" was then added thereto with stirring at 25°C until homogeneous. For aiding in dispersion of the thickener in the formulation, 4 g of carbomer available from B.F. Goodrich, Inc. under the tradename, "Carbopol Ultrez" were added to 10 g of triisopropyl citrate available from Phoenix Chemical Company under the tradename, "PELEMOL TIPC" in a separate beaker with hand stirring. Into the triisopropyl citrate mixture was then added 2 g. of methylparaben and 1 g of propylparaben with hand stirring until homogeneous to produce a pre-mixture. The pre-mixture was then added to the primary glass beaker with constant stirring until the resulting mixture was homogeneous.

Preparation of Final Composition:

- 2 g of triethanolamine available from Union Carbide under the tradename, "Trolamine 99%" was then added to the hydrophilic phase with constant stirring at 25°C until homogeneous. The resulting mixture was then added to the lipophilic phase at constant temperature with stirring until homogeneous. The final emulsion contains the components as set forth in Table F:

Table F: Emulsion Components

Chemical Name	Trade Name	%(wt/wt)
PEG-30 dipolyhydroxystearate	Arlacel P-135	1.1
Isononyl isononanoate	Wickenol	3.0
Hexyldecyl benzoate and butyloctyl benzoate	Hallstar AB	3.0
Cyclomethicone	Dow 344 Fluid	3.0
Vitamin A alcohol and Tween 20	Retinol 50C	0.69
Water	Water	86.320
Carbomer	Carbopol Ultrez	0.40
PEG-8 caprylic/capric glycerides	Trivasol BW	1.0
Methylparaben	Methylparaben	0.20
Propylparaben	Propylparaben	0.10
Triisopropyl citrate	Pelemol TIPC	1.0
NaOH	NaOH	0.190

Example 12 – Luminosity of the Formulation of Example 10

- Digital images of the right side and the left side of a Caucasian woman's face was taken using a digital camera available from Fujix (Model No.: DCS 505) equipped with a 60 mm macro lens under strobe light conditions at F8 and 1/125 seconds. The camera lens was filtered with a CG-395 filter, and the strobe light source was filtered with a combination of a UG-11 filter and a KG-5 filter. These images are illustrated in FIG. 3(a) and FIG. 3(b), respectively.
- After approximately 0.09 grams of the 0.3% retinol formulation prepared in Example 10 was applied to about a 20 cm² site on the suborbital (cheek) area of the right side and the

left side of the woman's face, digital images were taken thereof under the above conditions as illustrated in FIG. 3(c) and FIG. 3(d), respectively. Using PHOTOSHOP software available from Adobe Inc., the digital image of each site was analyzed for average pixel intensity or luminosity. Luminosity, as used herein, is an indication of brightness of a given area as measured on a scale of 1 to 255, wherein the latter is the most luminescent. Using the 0.3% retinol concentration value, the pixel intensity change, as determined by the difference in pixel intensity between the base surface and the treated surface, for both the treated right side and left side of the face was plotted as a function thereof as illustrated in FIG. 5 (a).

The formulation was then rinsed from the right side of the face, and a digital image was taken of the site under the above conditions as illustrated in FIG. 3(e). The formulation was then wiped twice using a Kimwipe tissue available from Kimberly Clark from the left side of the face, and a digital image was taken of the site under the above conditions as illustrated in FIG. 3(f). The pixel intensity change for the rinsed right side and the wiped left side was plotted on the graph of FIG. 5(a), then the respective deposited retinol concentrations were interpolated therefrom to be 0.145% and 0.1%, respectively.

This Example showed that the formulation of the present invention is not only a cleanser, but it also effectively deposits active agents, such as retinol, onto the skin. A significant amount of the agents remained on the skin after the formulation was removed therefrom. Moreover, this Example highlighted that when the cleanser composition of the present invention contains a 0.3% retinol active agent, it deposited the same amount of retinol on the skin as a leave-on product containing 0.145% retinol (when the compositions was removed via rinsing with water) and a leave-on product containing a 0.1% retinol (when the composition was removed via wiping).

Example 13 – Luminosity of the Formulation of Example 11

The procedure set forth in Example 12 was repeated using the formulation of Example 11 instead of that of Example 10. The pre-treatment images are illustrated in FIG. 4(a) and FIG. 4(b), respectively.

The formulation-containing images are illustrated in FIG. 4(c)(right side) and FIG. 4(d)(left side). Using the 0.3% retinol concentration value, the pixel intensity change for the treated right side and left side of the face was plotted as a function thereof as illustrated in FIG. 5 (d).

The digital image of the washed site is illustrated in FIG. 4(e), and the image of the wiped side is illustrated in FIG. 4(f). The pixel intensity change for the rinsed right side and the wiped left side was plotted on the graph of FIG. 5(b), then the respective deposited retinol concentrations were interpolated therefrom to be 0.135% and 0.072%, respectively.

This Example showed that the formulation of the present invention was not only a cleanser, but it also effectively deposited active agents, such as retinol, onto the skin. These agents remained present on the skin after the formulation was removed therefrom.

Moreover, this Example highlighted that when the cleanser composition of the present invention contained a 0.3% retinol active agent, the cleanser deposited the same amount of retinol on the skin as a leave-on product containing 0.135% retinol (when the composition was removed via rinsing with water) and a leave-on product containing a 0.072% retinol (when the composition was removed via wiping).

10 **Example 14: Preparation of Oil-in-Water Emulsion Containing DMAE**

Preparation of Lipophilic Phase:

10 g of steareth-2 available from Uniqema under the tradename "Brij 72", 8.5 g of isoceteth-20 also available from Uniqema under the tradename "Arlasolve 200", 10 g. of isononyl isononanoate available from Alzo, Inc., under the tradename "Wickenol 151", 10 g. of Isostearyl palmitate available from Brooks Industries under the tradename "Loronate OP", 10 g of cetyl octanoate, available from Brooks Industries, under the tradename "Loronate CIO," 10 g of pentaerythritol tetraoctanoate also available from Brooks Industries, under the tradename "Loronate PT," and 10 g of cyclomethicone available from Dow Coming under the tradename, "Dow 345 Fluid" were combined into a glass beaker at a temperature of 50 °C and stirred until homogeneous.

Preparation of Hydrophilic Phase:

602.5 g of deionized water were weighed into a primary glass beaker and heated to 78-82°C. With constant agitation, 4 g of PVM/MA Decadiene Crosspolymer available from ISP under the tradename, "Stabileze QM" was added thereto and held at 78-82°C until homogenous. This mixture was then cooled to 40-50°C, during which time, 1 g. of disodium EDTA, 10 g. of hexylene glycol, 7.5 g. of PEG-6 caprylic/capric glycerides available from Croda, Inc. under the tradename, "Glycerox 767", and 10 g. of PEG-150 pentaerythrityl tetrastearate also available from Croda under the tradename, "Crothix Liquid," were added to the primary beaker with constant stirring.

30 Preparation of Final Composition:

When both the lipophilic phase and the hydrophilic phase were at a temperature of 40°C -50°C, the lipophilic phase was added to the hydrophilic phase with constant stirring. In a separate beaker, 30 g. of 2-(dimethylamino) ethanol, available from BASF under the tradename DMAE, and 50 g. of L-tyrosine available from Ajinomoto under the tradename "L-Tyrosine" were added to 150 g. of water, and mixed until homogenous. This premix was then added to the primary beaker with constant stirring. 10 g of Nylon-12 available from Kobo Products, Inc., under the tradename "SP-10", 5 g of talc available from Luzenac America under the tradename, "Windsor Talc 66", 10 g of silicone quaternium-13 available from Biosil Industries under the tradename, "Biosil Basics SPQ," and 10 grams of a phenoxyethanol, methylparaben, butylparaben, ethylparaben and propylparaben solution available from Nipa under the tradename "Phenonip" were added separately to the primary beaker with constant stirring. The entire mixture was adjusted to a pH of 7.0-7.5 with a 70% aqueous solution of glycolic acid, and homogenized for 2 minutes at medium power with a Gifford-Wood homogenizer.

After about 1 ml to about 10 ml of the resulting formulation is applied to the facial skin of consumers, the consumers perceive that their facial skin appears and feels firmer and "lifted."

Example 15: *Preparation of Oil in Water Emulsion Containing a Polymeric Emulsifier and Colloidal Oat Flour*

Preparation of Hydrophilic Phase:

Into a primary glass beaker containing 850.70 g of deionized water, 10g of Colloidal Oat Flour available from Quaker was added thereto with stirring at 25°C until a homogeneous, smooth slurry was achieved. 2.5 g. of Acrylates/C10-30 Alkyl Acrylate Crosspolymer available from B.F. Goodrich, Inc. under the tradename, "Pemulen TR-1" and 2.5 g. of Carbomer, also available from B.F. Goodrich, Inc. under the tradename "Carbopol Ultrez" were then added to the primary beaker and mixed with slower agitation until homogenous. Into a separate beaker was added 7.5 g. of sucrose cocoate available from Croda, Inc. under the tradename, "Crodesta SL-40," 7.5 g. of PEG -6 Capric/caprylic glycerides available from Croda, Inc. under

the tradename, "Glycerox 767," 10 g of hexylene glycol, 3 g. of methylparaben and 0.5 g of propylparaben with hand stirring until homogeneous to produce a pre-mixture. The pre-mixture was then added to the primary glass beaker with constant stirring until the resulting mixture was homogeneous.

5 Preparation of Final Composition:

20 g of isostearyl palmitate, available from Brooks Industries, under the tradename "Loronate OP," 20 g of isononyl isononanoate, available from Alzo, Inc. under the tradename, "Wickenol 151," 20 g of cetyl octanoate, available from Brooks Industries, under the tradename "Loronate CIO," 20 g of pentaerythritol tetraoctanoate available from Brooks Industries, under
10 the tradename "Loronate PT," and 20 g of cyclomethicone available from Dow Corning under the tradename, "Dow 345 Fluid" were each added separately to the primary beaker with constant stirring at 25°C until homogeneous. 2.5 g of Tetrasodium EDTA and , 6.8 g. of a 20% aqueous solution of sodium hydroxide was then added thereto with stirring at a temperature of 25 °C. The resulting mixture was then mixed for 15 minutes.

15

**Example 16: Preparation of Oil in Water Emulsion
Containing a Polymeric Emulsifier and Colloidal Oat Flour**

20 Preparation of Preservative Pre-Blend

4 g of methylparaben, 1 g of propylparaben, 7.5 g of PEG-6 capric/caprylic glycerides available from Croda, Inc. under the tradename, "Glycerox 767," 7.5 g of sucrose cocoate also available from Croda, Inc. under the tradename, "Crodesta SL-40," and 10 g of hexylene glycol were combined with mixing under ambient conditions until homogeneous.

25

Preparation of Emulsion:

Into a primary glass beaker containing 852.5 g of Purified Water (USP), 10g of Colloidal Oat Flour available from Quaker were added thereto with stirring at about 200 rpm and a temperature of about 20 °C to about 30 °C until a homogeneous, smooth slurry was
30 achieved. 2.5 g. of Acrylates/C10-30 Alkyl Acrylate Crosspolymer available from B.F. Goodrich, Inc. under the tradename, "Pemulen TR-1" and 2.5 g. of Carbomer, also available

from B.F. Goodrich, Inc. under the tradename "Carbopol Ultrez" were then added thereto and mixed with slower agitation until homogenous. After adding the Preservative Pre-blend with increased mixing at about 200 rpm thereto, the following components were sequentially added thereto with constant stirring at about 20°C to about 30°C until homogeneous, with intervals of

5 5 minutes between the addition of each respective component: 20 g of isononyl isononanoate, available from Alzo, Inc. under the tradename, "Wickenol 151," 20 g of cyclomethicone available from Dow Corning under the tradename, "Dow 345 Fluid", 20 g of isostearyl palmitate, available from Brooks Industries, under the tradename "Loronate OP," 20 g of cetyl octanoate, available from Brooks Industries, under the tradename "Loronate CIO," 20 g of

10 pentaerythritol tetraoctanoate available from Brooks Industries, under the tradename "Loronate PT." 2.5 g of Tetrasodium EDTA and enough of a 20% aqueous solution of sodium hydroxide was then added thereto with stirring at a temperature of about 20°C to about 30°C to produce a final mixture having a pH of 5.9 to 6.5. The resulting mixture was then mixed until homogeneous.

15

Example 17— Consumer Testing of Formulation of Example 16 Formula

One hundred and twenty-five mothers of babies aged 24 months and younger used both the formulation prepared in accordance with Example 16 as well as a cleanser available from Galderma Laboratories, Inc. under the tradename, "CetaPhil Gentle Skin Cleanser."

20 The mothers used each product for a minimum of at least 3 to 7 times for a one week period. When using the product in a rinse-off fashion, the mothers first poured the product onto a moistened hand or wet cloth then applied the product to the desired location on the babies' skin. After rubbing the product gently on the skin, the product was rinsed therefrom with water. When using the product in a wipe-off fashion, the mothers applied a liberal amount to the

25 desired location on the babies' skin and rubbed gently. The excess product was then removed therefrom with a soft cloth or tissue.

The results of the study are shown in Table G below:

Table G Comparative Study of Example 16 Formulation to Cetaphil

Characteristic that either Completely describes or very well describes the product at issue	Formulation of Example 16	Cetaphil
1) Good for Sensitive Skin	87*	82
2) Nonirritating to Skin	92	91
3) Won't Dry or Irritate the Most Sensitive Skin	89	78
4) Cleanses Extra Gently	87	86
5) Good for Everyday or Regular Use	94	84
6) Cleans Without Drying Skin	92	85
7) Relieves Baby's Dry Skin	74	65
8) Can Be Use With or Without Water	93	83
9) Effectiveness as a Cleanser	82	76
10) Good for Use All Over the Body	90	85
11) Makes Baby's Skin Feel Soft and Smooth	89	74
12) Helps Baby's Skin Look and Feel Healthier	70	66
13) Helps Baby's Skin Retain its Natural Moisture	82	72
14) Relieves Baby's Itchy Skin	51	46
15) Leaves Baby's Skin Feeling Clean	86	75
16) Won't Sting or Irritate Eyes	54	54

*These numbers indicate the percentage of the mothers that indicated that the identified product possessed the given characteristic.

- This Example showed that the cleanser of Example 16 significantly outperformed the commercial product with respect to the majority of characteristics set forth in Table G.

Example 18 – Preparation of Oil-in-Water Emulsion with Non-Ionic Emulsifier

Preparation of Lipophilic Phase:

- 20 g of isostearyl palmitate, available from Brooks Industries, under the tradename "Loronate OP," 20 g of isononyl isononanoate, available from Alzo, Inc. under the tradename, "Wickenol 151," 20 g of cetyl octanoate, available from Brooks Industries, under the tradename

"Loronate CIO," 20 g of pentaerythritol tetraoctanoate available from Brooks Industries, under the tradename "Loronate PT," were combined into a glass beaker at a temperature of 25 °C and stirred until homogeneous.

Preparation of Hydrophilic Phase:

- 5 Into a primary glass beaker containing 859.7 g of deionized water, 2 g of carbomer available from B.F. Goodrich, Inc. under the tradename, "Carbopol ETD 2020" , and 1 g of C10-C30 alkyl acrylate/crosspolymer commercially available from B.F. Goodrich under the tradename, "Pemulen TR1" were added thereto with stirring at a temperature of 25 °C until dispersed. While heating the mixture to 75 °C, 1.2 g of tromethamine, 1g of EDTA, 7.5 g. of
- 10 PEG -6 Capric/caprylic glycerides available from Croda, Inc. under the tradename, "Glycerox 767," 10 g of hexylene glycol, 4 g. of methylparaben and 1 g of propylparaben were added with constant stirring until the resulting mixture was homogeneous. After the mixture reached a temperature of 75 °C, 10 g. of a mixture of sorbitan stearate and sucrose cocoate available from Uniqema under the tradename, "Arlatone 2121," were added thereto with stirring for 30
- 15 minutes at constant temperature.

Preparation of Final Composition:

- After the lipophilic phase was heated to a temperature of 75 °C, it was then added to the hydrophilic phase with constant stirring at 75°C until homogeneous. After the mixture was then cooled to 35 °C, 20 g of cyclomethicone available from Dow Corning under the
- 20 tradename, "Dow 345 Fluid" was added thereto. After the mixture was cooled to 25 °C, 0.4 g. of tromethamine was then added thereto with stirring at constant temperature such that the resulting mixture had a pH of 5.5.

Example 19: Comparison of Make-Up Removability

- 25 A waterproof mascara available from Gemey under the tradename "Waterproof Gemey Noir," is copiously coated on a 16 cm² area on a forearm. This area is then wiped four times with a cotton ball soaked with 2 ml of the formula of Example 2. Visual assessment of

the area is noted. This procedure is repeated on the same area and a second visual assessment is noted. This procedure is repeated on the same area and a third visual assessment is noted.

This procedure is repeated on an alternative areas of the forearm, but wherein the cleanser of Example 2 is replaced with the cleanser of Example 18 and the cleanser available from Johnson's under the tradename, "Johnson's pH 5.5 3-in-1 Cleansing Lotion," respectively.

Although all three cleansers effectively remove the mascara, this Example shows that the cleansers of Example 2 and Example 18 remove mascara more quickly than the commercial cleanser. In addition, all three cleansers are aesthetically acceptable according to consumer standards.

Example 20: Comparison of Make-Up Removability of Impregnated Wipes

The formulation of Example 18 is prepared, but wherein: 1) the carbomer and the C10-C30 alkyl acrylate/crosspolymer were replaced with 1 g of xanthan gum; 2) and 30 g of C14 – 22 alkyl alcohol and C12 –C20 alkyl glucoside emulsifier available from Seppic under the tradename "Montanov L" was also added to the lipophilic phase; and 3) and 4 g of C13-C14 isoparaffin/isostearyl isostearate/Na polyacrylate/polyacrylamide/polysorbate 60 available from Seppic under the tradename, "Sepigel 502," was added to the finished mixture as the final product was cooled to 25 C.

The emulsion was pumped on to a stack of 25 folded, uncoated spunlaced wipes comprised of a blend of about 35% polyester and 65% rayon in an amount equivalent to 325% of the total noncoated wipe weight.

The resulting wipes were then compared with Johnson pH5.5 3-in-1 Cleansing wipes and Pond's Cleansing Towelettes available from Unilever using the test method set forth in Example 19 wherein the cotton ball was replaced by the respective wipe.

This Example showed that the wipes prepared in accordance with this Example are more effective in removing waterproof mascara than the two commercial products as determined after the first, second, and third assessment. More specifically, the wipes of this

Example were significantly superior to the Pond's wipes with respect to mascara removal as determined after the third visual assessment.

Example 21: Preparation of Cleansing Composition

5

20 g of cyclomethicone available from the Dow Corning Corporation under the tradename, "DOW CORNING 345," 15 g of hexylene glycol, and 65 g of a mixture of hexyl decyl benzoate and butyl octyl benzoate available from the C.P. Hall Company under the tradename, "Hallstar AB" are sequentially added to a vessel with mixing at about 100 rpm under ambient conditions until the final mixture is homogeneous.

10

The resulting cleansing composition is effective in removing debris such as makeup from the skin.

We claim:

1. A cleansing composition comprising
 - a. a liquid silicone;
 - 5 b. a water dispersible component and
 - c. an ester.
2. The composition of claim 1 wherein the composition is comprised of, based upon the total weight of the composition,
 - a. from about 10 percent to about 35 percent of the liquid silicone;
 - 10 b. from about 10 percent to about 35 percent of the water dispersible component;
 - and
 - c. from about 55 percent to about 65 percent of the ester.
3. The cleansing composition of claim 1 wherein the silicone is selected from the group consisting of hexamethylsiloxane, dimethicone, dimethiconol, cyclomethicone, and
15 mixtures thereof.
4. The cleansing composition of claim 3 wherein the cyclomethicone is selected from the group consisting of cyclo tetradimethyl siloxane; cyclopentadimethyl siloxane, cyclohexadimethyl siloxane, cycloheptadimethyl siloxane, and mixtures thereof.
5. The cleansing composition of claim 1 wherein the water dispersible component is
20 selected from the group consisting of polyethylene glycol 400, hexylene glycol, propylene glycol, polypropylene glycol-10 methylglucose ether, ethoxydiglycol, polyethylene glycol-6 caprylic/capric glycerides, ethylene glycol monobutyl ether, triisopropyl citrate, polyethylene glycol-8 caprylic/capric glycerides, 3-methoxy-3-methyl-1-butanol, dimethyl isosorbide, polyethylene-6 caprylic/capric triglyceride, and mixtures thereof.
6. The cleansing composition of claim 5 wherein the water dispersible component is
25 selected from the group consisting of hexylene glycol, dimethyl isosorbide, polyethylene glycol-6 caprylic/capric glyceride, and mixtures thereof.
7. The cleansing composition of claim 1 wherein the water dispersible component is comprised of, based upon the total weight percent of the cleaning composition,
30 a) from about 5 percent to about 15 percent of hexylene glycol;
- b) from about 5 percent to about 10 percent of polyethylene-6 caprylic/capric triglyceride.

8. The cleansing composition of claim 1 wherein the ester is selected from liquid esters that either possess a structural means for ensuring the liquidity of the ester or are heterogeneous in nature.

5 9. The cleansing composition of claim 1 wherein the ester is selected from the group consisting of

a) a branched C₅ to C₂₂ alkyl alcohol ester of an aromatic acid;

b) a straight-chained or branched C₅ to C₂₂ alkyl acid esters of optionally ethoxylated/propoxylated polyols having from about 3 carbon atoms to about 7 carbon atoms;

10 c) branched C₅ to C₂₂ alkyl alcohol esters of branched polyacids;

d) branched or straight-chained C₅ to C₂₂ alkyl acid esters of branched and/or unsaturated C₅ to C₂₂ alkyl alcohols;

e) branched or unsaturated C₅ to C₂₂ alkyl alcohol esters of an acid selected from the group consisting of adipic acid, succinic acid, sebacic acid, maleic acid, and mixtures thereof

15 f) polyether interrupted fatty acid esters;

g) benzoic acid ester of heterogeneous alcohols having from about 8 carbon atoms to about 22 carbon atoms; and

h) mixtures thereof,

20 10. The cleansing composition of claim 9 wherein the ester is selected from the group consisting of straight-chained or branched C₅ to C₂₂ alkyl acid esters of optionally ethoxylated/propoxylated polyols; benzoic acid esters of heterogeneous alcohols; and mixtures thereof.

25 11. The cleansing composition of claim 9 wherein the ester is selected from the group consisting of butyloctyl salicylate; hexyldecyl benzoate; and butyloctyl benzoate; alkyl benzoates having from about 12 carbon atoms to about 15 carbon atoms; and mixtures thereof.

12. The cleansing composition of claim 11 wherein the ester is selected from the group consisting of hexyldecyl benzoate, butyloctyl benzoate, and mixtures thereof.

30 13. The cleansing composition of claim 9 wherein the ester is selected from the group consisting of pentaerythritol tetraoctanoate; trimethylolpropane trioctanoate; trioctanoin; pentaerythrityl tetrapelargonate; sorbitan trioleate; caprylic/capric triglyceride; neopentyl alcohol tetraoctanoate, and mixtures thereof.

14. The cleansing composition of claim 13 wherein the ester is selected from the group consisting of caprylic/capric triglyceride; pentaerythritol tetraoctanoate; trimethylolpropane trioctanoate; pentaerythrityl tetrapelargonate; and mixtures thereof.
- 5 15. The cleansing composition of claim 9 wherein the ester is selected from the group consisting of branched alkyl alcohol esters of branched polyacids, wherein the alkyl alcohol is optionally substituted and contains from about 3 carbon atoms to about 22 carbon atoms.
16. The cleansing composition of claim 15 wherein the ester is trioctylododecyl citrate and mixtures thereof.
- 10 17. The cleansing composition of claim 9 wherein the ester is selected from the group consisting of tridecyl neopentanoate, isostearyl palmitate, cetyl ricinoleate, cetyl octanoate, isononyl isononanoate, butyl stearate, octyldodecyl soyate, tridecyl erucate, octyldodecyl erucate/eicosil erucate, and mixtures thereof.
- 15 18. The cleansing composition of claim 17 wherein the ester is selected from the group consisting of cetyl octanoate, isostearyl palmitate, isononyl isononanoate, and mixtures thereof.
19. The cleansing composition of claim 9 wherein the ester is selected from the group consisting of diisopropyl adipate, dioctyl sebacate, dioctyl succinate, dioctyl maleate, diisostearyl adipate, diethyl sebacate, and mixtures thereof.
- 20 20. The cleansing composition of claim 19 wherein the ester is selected from the group consisting of diethyl sebacate, dioctyl sebacate, diisostearyl adipate, and mixtures thereof.
- 25 21. The cleansing composition of claim 9 wherein the ester is selected from the group consisting of laureth-2 benzoate; C_8 to C_{22} fatty alkyl (optionally polypropylenoxy) polyethyleneoxy carboxylate esters derived from an alcohol having from about 1 carbon atom to about 22 carbon atoms; and mixtures thereof.
22. The cleansing composition of claim 21 wherein the ester is isopropyl propylene glycol-2-isodeceth-7 carboxylate.
- 30 23. The cleansing composition of claim 9 wherein the ester is selected from at least two of the following esters:
- a) branched C_5 to C_{22} alkyl alcohol esters of an aromatic acid;
 - b) branched or straight-chained C_5 to C_{22} alkyl acid esters of branched or unsaturated C_5 to C_{22} alkyl alcohols; and

b) branched or straight-chained C₅ to C₂₂ alkyl acid esters of branched or unsaturated C₅ to C₂₂ alkyl alcohols; and

5 c) straight-chained or branched C₅ to C₂₂ alkyl acid esters of optionally ethoxylated/propoxylated polyols.

24. The cleansing composition of claim 9 wherein the ester is a mixture of, based upon the total weight percent of the esters,:

a) from about 30 percent to about 80 percent of branched or straight-chained C₅ to C₂₂ alkyl acid esters of branched or unsaturated C₅ to C₂₂ alkyl alcohols;

10 b) from about 10 percent to about 50 percent of branched C₅ to C₂₂ alkyl alcohol esters of an aromatic acid; and

c) from about 10 percent to about 50 percent of straight-chained or branched C₅ to C₂₂ alkyl acid esters of optionally ethoxylated/propoxylated polyols.

15 25. The cleansing composition of claim 9 wherein the ester is a mixture comprised of, based upon the total weight percent of the ester,:

a) from about 15 percent to about 50 percent isononyl isononanoate;

b) from about 15 percent to about 50 percent isostearyl palmitate;

c) from about 15 percent to about 50 percent cetyl octanoate; and

d) from about 15 percent to about 50 percent pentaerthritol tetraoctanoate.

20 26. A cleaning system comprising:

a. the cleansing system of claim 1;

b. water; and

c. a polymeric emulsifier and/or a thickener.

25 27. The cleansing system of claim 26 comprising, based upon the total weight of the cleansing system:

a. at least 5 percent of the cleansing system of claim 1;

b. from about 70 percent to about 98 percent of water; and

c. from about 0.5 to about 1.5 percent of a polymeric emulsifier and/or thickener.

30

28. The cleansing system of claim 26 wherein the polymeric emulsifier is polyethylene glycol-30 dipolyhydroxystearate; dimethicone copolyol; substituted acrylates; and mixtures thereof.

5 29. The cleansing system of claim 26 wherein the thickener is selected from the group consisting of carbomers, acrylate copolymers, hydroxyethylcellulose modified with cetyl ether groups, polyvinylmethyl ether/maleic anhydride (PVM/MA) decadiene crosspolymer, and mixtures thereof.

10 30. The cleansing system of claim 26 wherein the thickener is selected from the group consisting of acrylates/aminoacrylates copolymer, acrylates/steareth-20 methacrylate copolymer; acrylates/ceteth-20 itaconate copolymer, acrylates/steareth-20 itaconate copolymer, carbomers, modified hydroxycellulose, polyvinylacetate/maleic anhydride (PVA/MA) decadiene crosspolymer, and mixtures thereof.

15 31. The cleansing system of claim 26 further comprising, based upon the total weight of the cleansing system, from about 1 percent to about 3 percent of a cleansing enhancer.

 32. The cleansing system of claim 31 wherein the cleansing enhancer is a nonfoaming surfactant and/or a non-ionic emulsifier.

20 33. The cleansing system of claim 32 wherein the nonfoaming surfactant is selected from the group consisting of sucrose cocoate, sucrose stearate and mixtures thereof.

 34. The cleansing system of claim 32 wherein the non-ionic emulsifier is selected from the group consisting of isoceteth 20, oleth-2, mixture of PEG-40 hydrogenated castor oil and trideceth-9, Poloxamer 184, laureth-4, sorbitan trioleate, polyoxyethylene-(2) oleyl ether, sorbitan stearate, cetearyl glucoside, glyceryl oleate, and mixtures thereof.

25 35. The system of claim 26 further comprising a benefit agent.

 36. The system of claim 35 wherein the benefit agent is selected from the group consisting of vasoconstrictors, collagen enhancers, anti-edema agents, depigmentation agents; reflectants; detangling/wet combing agents; film forming polymers; humectants; amino acid agents; antimicrobial agents; allergy inhibitors; anti-acne agents; anti-aging agents; anti-wrinkling agents, antiseptics; analgesics; antitussives; antipruritics; local anesthetics; anti-hair loss agents; hair growth promoting agents; hair growth inhibitor agents; antihistamines; antiinfectives; inflammation inhibitors; anti-emetics; anticholinergics; vasoconstrictors; vasodilators; wound healing promoters; peptides, polypeptides and proteins; deodorants and anti-perspirants; medicament agents; skin
30 mollients and skin moisturizers; skin firming agents, hair conditioners; hair softeners; hair
35 moisturizers;

vitamins; tanning agents; skin lightening agents; antifungals; depilating agents; shaving preparations; external analgesics; perfumes; counterirritants; hemorrhoidals; insecticides; poison ivy products; poison oak products; burn products; anti-diaper rash agents; prickly
 5 heat agents; make-up preparations; vitamins; amino acids and their derivatives; herbal extracts; retinoids; flavenoids; sensates; anti-oxidants; skin conditioners; hair lighteners; chelating agents; cell turnover enhancers; coloring agents; pigments; sunscreens and mixtures thereof.

37. The system of claim 35 wherein the benefit agent is selected from the group
 10 consisting of feverfew, centella asiatica, olive leaf, wheat protein, oat oil, lycopene, DMAE, soy and derivatives thereof, colloidal oatmeal, sulfonated shale oil, elubiol, 6-(1-piperidinyl)-2,4-pyrimidinediamine-3-oxide, finasteride, ketoconazole, salicylic acid, zinc pyrithione, coal tar, benzoyl peroxide, selenium sulfide, hydrocortisone, sulfur, menthol, pramoxine hydrochloride, tricetylammonium chloride, polyquaternium 10, panthenol, panthenol
 15 triacetate, vitamin A and derivatives thereof, vitamin B and derivatives thereof, vitamin C and derivatives thereof, vitamin D and derivatives thereof, vitamin E and derivatives thereof, vitamin K and derivatives thereof, keratin, lysine, arginine, hydrolyzed wheat proteins, hydrolyzed silk proteins, octyl methoxycinnamate, oxybenzone, minoxidil, titanium dioxide, zinc dioxide, retinol, erythromycin, tretinoin, and mixtures thereof.

38. The system of claim 34 wherein the benefit agent is present in an amount, based upon the total weight of the system, from about 0.001 percent to about 5.0 percent.

39. The system of claim 26 further comprised of, based upon the total weight of the system, from about 5 percent to about 15 percent of a foaming surfactant.

40. A method of treating hair loss comprising topically applying the system of claim
 25 25 with an effective amount of a hair loss treatment agent to a desired location on an animal or human.

41. The method of claim 40 wherein the hair loss treatment agent is selected from the group consisting of minoxidil, 6-(1-piperidinyl)-2,4-pyrimidinediamine-3-oxide, N'-cyano-N-(tert-pentyl)-N'-3-pyridinyl-guanidine, finasteride, retinoids and derivatives thereof,
 30 ketoconazole, elubiol or mixtures thereof.

42. A method for inhibiting hair growth comprising topically applying the system of claim 26 with a hair growth inhibiting agent to a desired area on an animal or human for inhibiting hair growth.

43. The method of claim 42 wherein the benefit agent is selected from the group
 35 consisting of serine proteases, retinol, isotretinoin, betamethoisone, alpha-tocophenol and derivatives thereof, and mixtures thereof.

5 44. A method for treating acne comprising topically applying a mixture of the system of claim 26 and an effective amount of an anti-acne agent to the skin of an animal or human at a desired area.

45. The method of claim 44 wherein the anti-acne agent is selected from the group consisting of benzoyl peroxide, retinol, elubiol, antibiotics, salicylic acid, and mixtures thereof.

10 46. A method for reducing the signs of aging and other manifestations of photodamage comprising topically applying a mixture of the system of claim 26 and an effective amount of an anti-aging agent to the skin of an animal or human at a desired area.

47. The method of claim 46 wherein the anti-aging agent is selected from the group consisting of retinoids, anti-oxidants, alpha-hydroxy acids, beta-hydroxy acids and mixtures thereof.

15 48. A method for depigmenting the skin comprising topically applying the system of claim 26 and an effective amount of a depigmentation benefit agent to the skin of an animal or human at a desired area.

49. The method of claim 48 wherein the depigmentation agent is selected from the group consisting of retinol, Kojic acid, hydroquinone, and mixtures thereof.

20 50. A method for treating the symptoms and/or the diseases of dandruff, seborrheic dermatitis and/or psoriasis, comprising topically applying a mixture of the system of claim 26 and an effective amount of a benefit agent capable of treating the symptoms to the skin of an animal or human at a desired area.

25 51. The method of claim 50 wherein the benefit agent is selected from the group consisting of shale oil and derivatives thereof, elubiol, ketoconazole, coal tar, salicylic acid, zinc pyrithione, selenium sulfide, hydrocortisone, sulfur, menthol, pramoxine hydrochloride, and mixtures thereof.

52. The use of the composition of claim 26 in a personal care product.

30 53. The composition of claim 52, wherein the personal care product is in the form of a gel, a bath, a wash, a mousse, a shampoo, a rinse, a lotion, a cream, a wipe, a brush, a sponge, or a spray.

54. The use of the composition of claim 26 as a make-up remover.

55. The composition of claim 26 in the form of an oil-in-water emulsion.

35 56. A foaming composition comprising:
a. a water dispersible component;
b. an ester;
c. water; and a

d. foaming surfactant.

57. The composition of claim 56 wherein the foaming surfactant has a column height of greater than about 20 mm as determined by the Miles-Ross Test and is selected from the group consisting of non-ionic surfactants, cationic surfactants, amphoteric surfactants, anionic surfactant, and mixtures thereof.

58. The composition of claim 57 further comprising a liquid silicone.

59. The composition of claim 57 further comprising at least one of the following:

- a) polymeric emulsifier and/or a thickener;
- b) a benefit agent; or
- c) a nonionic emulsifier.

60. A method for making an oil-in water emulsion comprised of:

a) neutralizing a hydrophilic thickening agent in a hydrophilic phase comprised of a polymeric emulsifier with an effective amount of a neutralizer under conditions sufficient after a lipophilic phase was combined with the hydrophilic phase.

61. The method of claim 60 wherein the hydrophilic thickening agent is selected from the group consisting of carbomers, acrylate copolymers, modified hydroxycellulose, polyvinylacetate/maleic anhydride (PVA/MA) decadiene crosspolymer, and mixtures thereof.

62. The method of claim 61 wherein the acrylate copolymers are selected from the group consisting of acrylates/aminoacrylates copolymer, acrylates/steareth-20 itaconate copolymer, acrylates/cetheth-20 itaconate copolymer, acrylates/steareth-20 methacrylate copolymer, and mixtures thereof.

63. The method of claim 59 wherein the hydrophilic phase is comprised of one or more of the following components: water, thickener, cleansing enhancer, nonfoaming surfactant, or water dispersible component.

64. The method of claim 60 wherein the lipophilic phase is comprised of one or more of the following components: silicone, ester, or polymeric emulsifier.

65. A method for making a water-in-oil emulsion comprised of:

a) neutralizing a hydrophilic thickening agent in a hydrophilic phase comprised of a polymeric emulsifier with an effective amount of a neutralizer under conditions sufficient before combining a lipophilic phase with the hydrophilic phase.

66. The method of claim 65 wherein the hydrophilic thickening agent is selected from the group consisting of carbomers, acrylate copolymers, modified hydroxycellulose,

polyvinyl methyl ether/maleic anhydride (PVM/MA) decadiene crosspolymer, and mixtures thereof.

67. The method of claim 65 wherein the acrylate copolymers are selected from the group consisting of acrylates/aminoacrylates copolymer, acrylates/steareth-20 itaconate copolymer, acrylates/ceteth-20 itaconate copolymer, acrylates/steareth-20 methacrylate copolymer, and mixtures thereof.

68. The method of claim 65 wherein the hydrophilic phase is comprised of one or more of the following: water, thickener, cleansing enhancer, nonfoaming surfactant, and water dispersible component.

69. The method of claim 65 wherein the lipophilic phase is comprised of one or more of the following: silicone, ester, and polymeric emulsifier.

70. A method for depositing a benefit agent into and/or onto the skin, hair and/or nails comprising applying a composition comprising:

- a. an optional liquid silicone;
- b. a water dispersible component;
- c. an ester;
- d. a polymeric emulsifier and/or a thickener; and
- e. an effective amount of a benefit agent

to a desired location on a human or animal.

71. The method of claim 70 wherein the composition is further comprised of a foaming surfactant, a cleansing enhancer, or a mixture thereof.

72. The method of claim 70 wherein the benefit agent is selected from the group consisting of colloidal oatmeal, olive leaf, soy and derivatives thereof, sulfonated shale oil, elubiol, 6-(1-piperidiny)-2,4-pyrimidinediamine-3-oxide, finasteride, ketoconazole, salicylic acid, zinc pyrithione, coal tar, benzoyl peroxide, selenium sulfide, hydrocortisone, sulfur, menthol, pramoxine hydrochloride, tricetylammonium chloride, polyquaternium 10, panthenol, panthenol triacetate, vitamin A and derivatives thereof, vitamin B and derivatives thereof, vitamin C and derivatives thereof, vitamin D and derivatives thereof, vitamin E and derivatives thereof, vitamin K and derivatives thereof, keratin, lysine, arginine, hydrolyzed wheat proteins, hydrolyzed silk proteins, octyl methoxycinnamate, oxybenzone, minoxidil, titanium dioxide, zinc dioxide, retinol, erthromycin, tretinoin, DMAE, and mixtures thereof.

73. A method of cleansing hair or skin comprised of applying an effective amount of the composition of claim 1 to a desired location.

74. A method of cleansing hair or skin comprised of applying an effective amount of the system of claim 26 to a desired location.

5 75. The cleansing composition of claim 26 wherein the water dispersible component is comprised of, based upon the total weight percent of the cleaning system,

- a) from about 0.1 percent to about 5 percent of hexylene glycol;
- b) from about 0.5 percent to about 3.0 percent of polyoxyethylene-6 caprylic/capric triglyceride.

10 76. A method for depositing a benefit agent into and/or onto the skin, hair and/or nails comprising applying a composition comprising:

- a) an optional liquid silicone;
- b) a water dispersible component
- c) an ester;
- 15 d) water;
- e) a foaming surfactant; and
- f) an effective amount of a benefit agent

to a desired location on a human or animal.

77. A cleaning composition comprised of

- 20
- a) a cyclomethicone liquid silicone;
 - b) a water dispersible component selected from the group consisting of hexylene glycol, PEG-6 caprylic/capric triglycerides, and mixtures thereof; and
 - c) an ester selected from the group consisting of isononyl isononanoate, isostearyl palmitate, cetyl octanoate, pentaerthritol tetraoctanoate, and mixtures thereof.

25 78. A cleaning system comprised of

- a) a cyclomethicone liquid silicone;
- b) a water dispersible component selected from the group consisting of hexylene glycol, PEG-6 caprylic/capric triglycerides, and mixtures thereof;
- c) an ester selected from the group consisting of isononyl isononanoate, isostearyl
- 30 palmitate, cetyl octanoate, pentaerthritol tetraoctanoate, and mixtures thereof;
- d) water; and

5 e) a PEG-30 dipolyhydroxystearate polymeric emulsifier and/or a carbomer thickener.

79. A foaming composition comprised of

a) water;

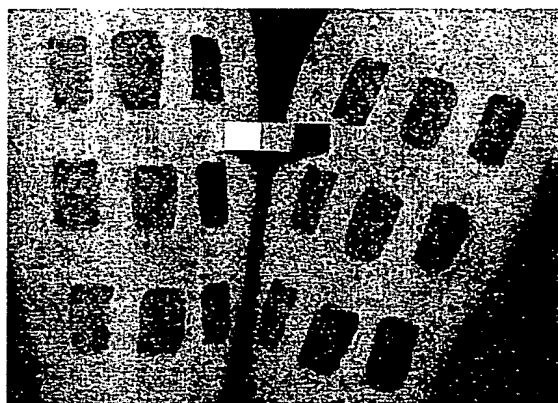
b) a water dispersible component selected from the group consisting of hexylene glycol, PEG-6 caprylic/capric triglycerides, and mixtures thereof;

10 c) an ester selected from the group consisting of isononyl isononanoate, isostearyl palmitate, cetyl octanoate, pentaerthritol tetraoctanoate, and mixtures thereof; and

d) a foaming surfactant selected from the group consisting of cocamide MEA, lauryl glucoside, PEG-50 tallow amide, cocamidopropylamine oxide, and mixtures thereof.

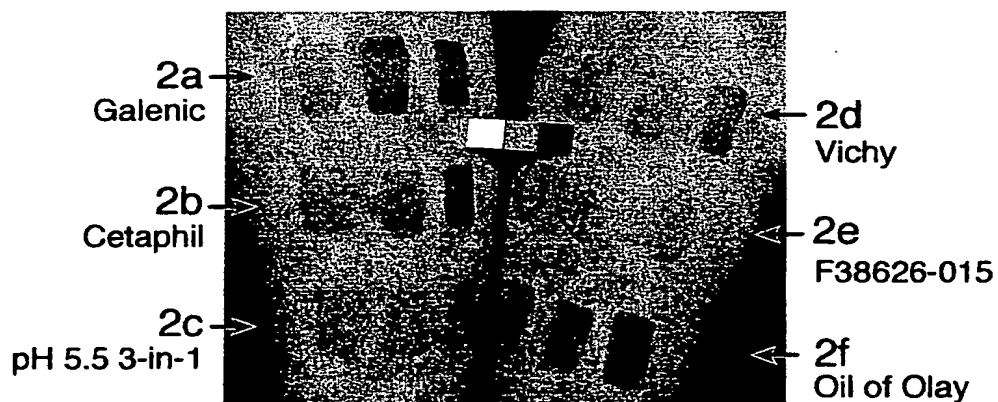
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FIG. 1



Before Removal

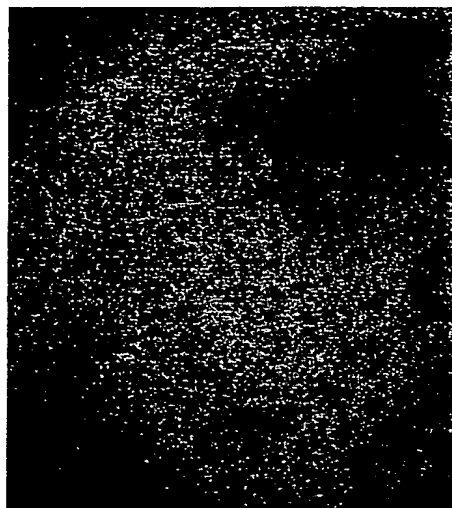
FIG. 2



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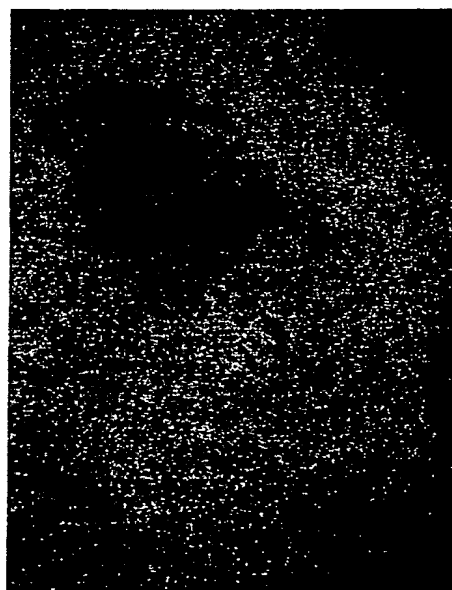
Retinol Fluorescence 0.3% rol Cleanser

FIG. 3(a)



Retinol Fluorescence 0.3% rol Cleanser

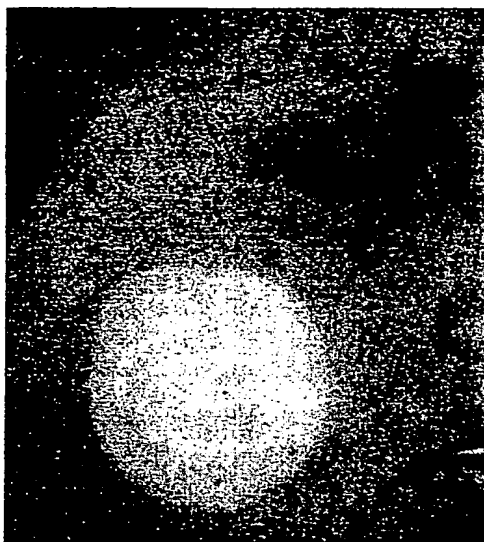
FIG. 3(b)



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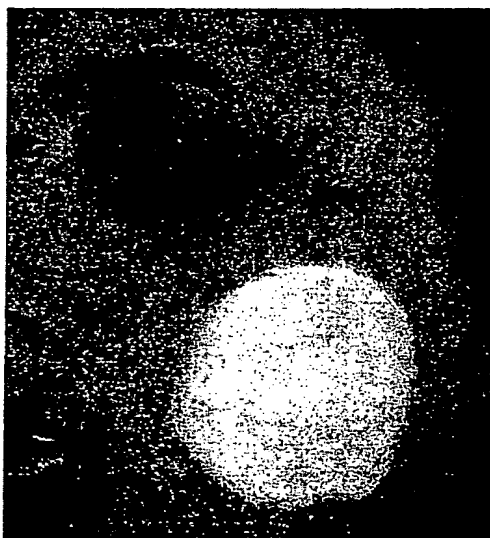
Luminosity Change = 78.84

FIG. 3(c)



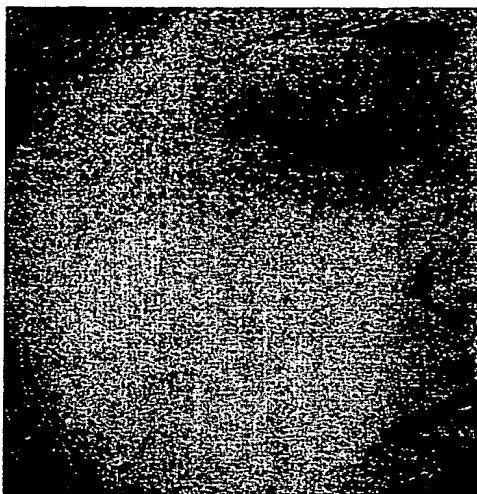
Luminosity Change = 82.33

FIG. 3(d)



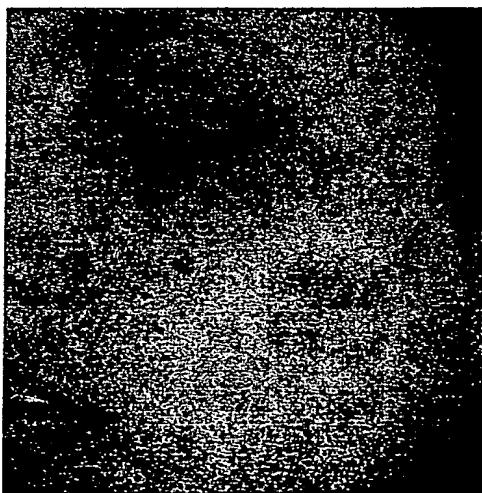
Rinse off Luminosity Change =37.86

FIG. 3(e)



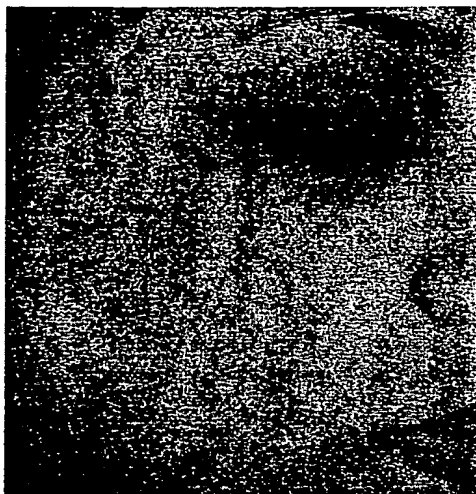
Wiped off Luminosity Change =27.19

FIG. 3(f)



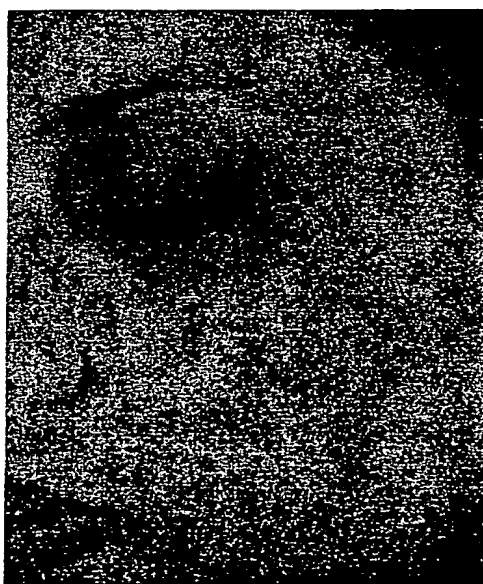
Retinol Fluorescence 0.3% rol Cleanser

FIG. 4(a)



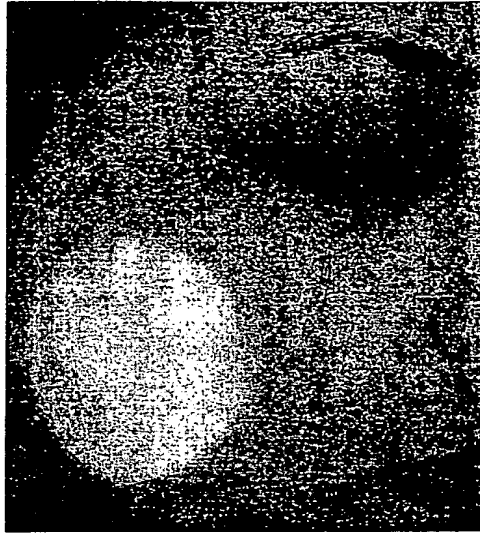
Retinol Fluorescence 0.3% rol Cleanser

FIG. 4(b)



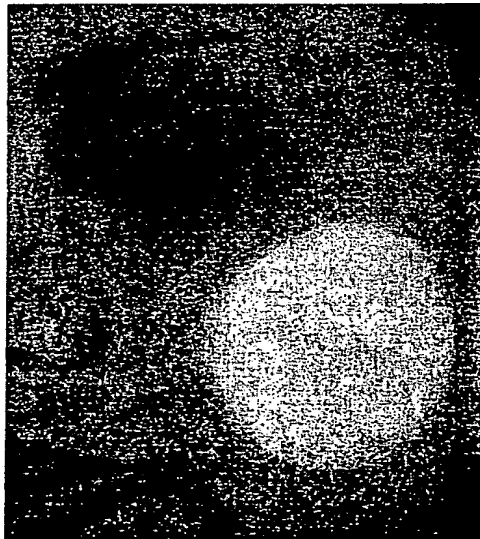
Luminosity Change = 65.92

FIG. 4(c)



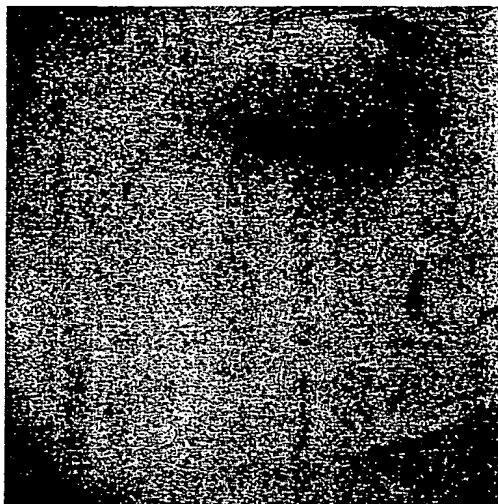
Luminosity Change = 70.42

FIG. 4(d)



Rinse off Luminosity Change =28.97

FIG. 4(e)



Wiped off Luminosity Change =17.01

FIG. 4(f)

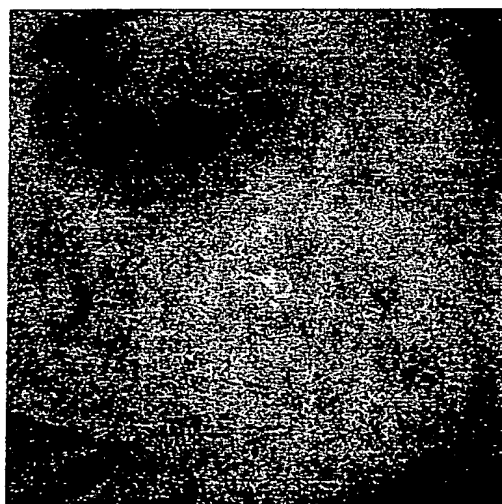


FIG. 5(a)

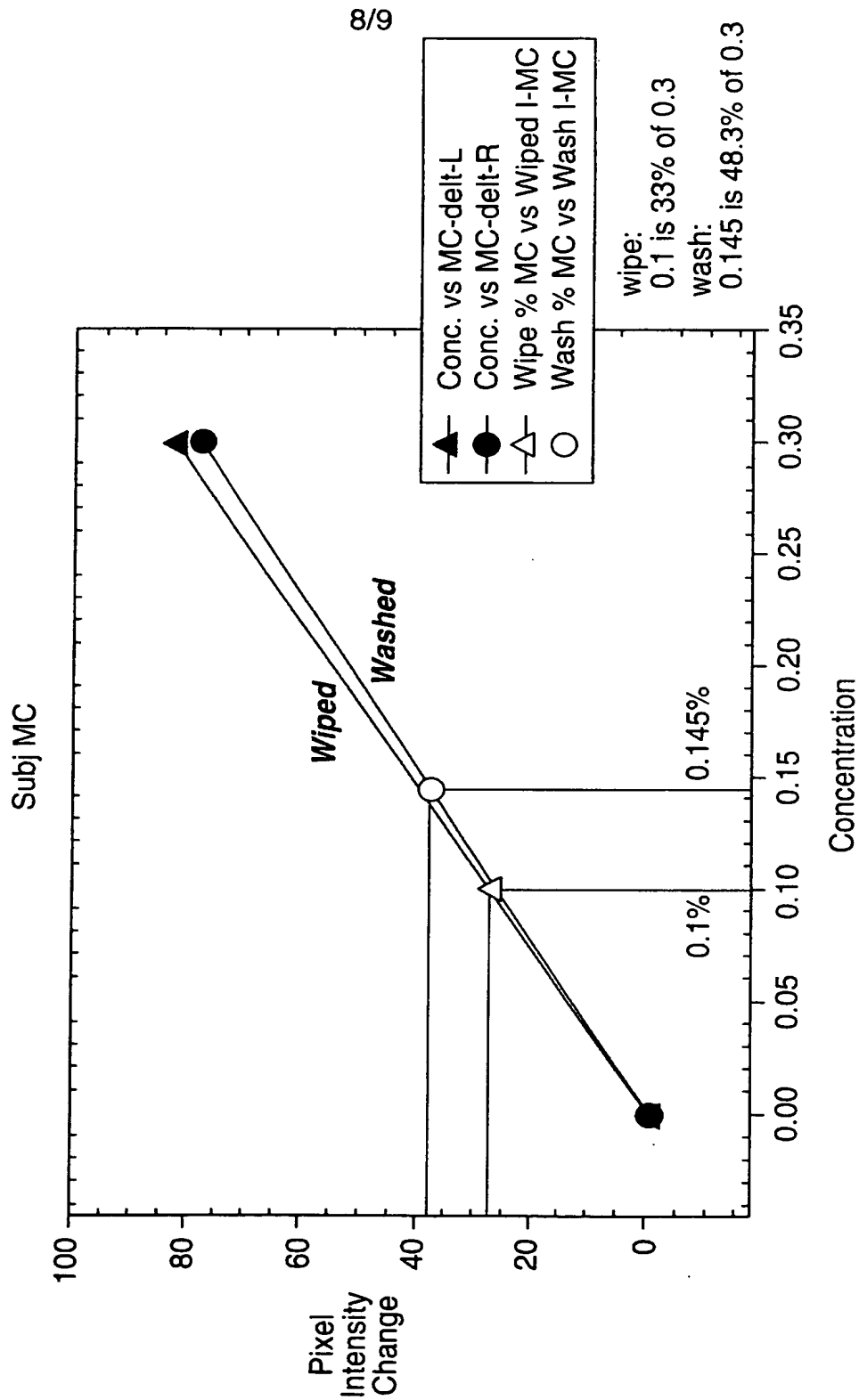
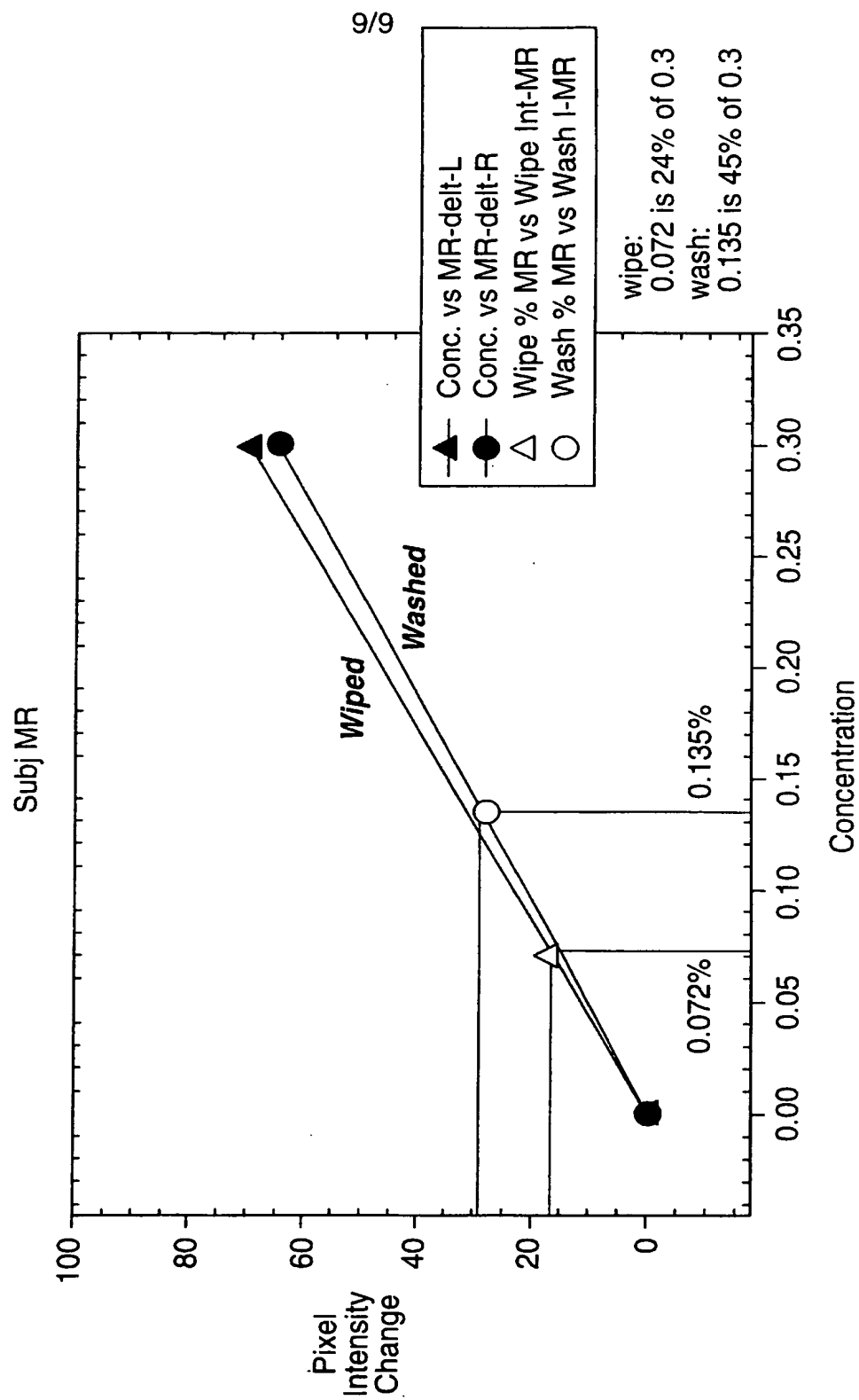


FIG. 5(b)



INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 00/17431

A. CLASSIFICATION OF SUBJECT MATTER IPC 7 A61K7/48 A61K7/50 A61K7/02		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC 7 A61K		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used) WPI Data, PAJ, EPO-Internal		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 95 05160 A (UNILEVER PLC ; UNILEVER NV (NL)) 23 February 1995 (1995-02-23) claims 1,5; example 8 ---	1-26, 55-78
A	WO 98 35652 A (EICKEN ULRICH ; JUNGO SYBILLE (CH); WELLA AG (DE); KISCHKA KARL HEI) 20 August 1998 (1998-08-20) claims 1-3,8-10 ---	1-33, 70-78
A	US 5 888 951 A (SIMON PASCAL ET AL) 30 March 1999 (1999-03-30) claim 1; example 2 ---	1-27, 55-58
A	WO 95 05145 A (UNILEVER PLC ; UNILEVER NV (NL)) 23 February 1995 (1995-02-23) page 9, line 37; claims 1,6 -----	1-26, 51-58, 72,73
<input type="checkbox"/> Further documents are listed in the continuation of box C. <input checked="" type="checkbox"/> Patent family members are listed in annex.		
* Special categories of cited documents :		
<div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 45%;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"&" document member of the same patent family</p> </div> </div>		
Date of the actual completion of the international search <div style="text-align: center; font-weight: bold;">6 November 2000</div>		Date of mailing of the international search report <div style="text-align: center; font-weight: bold;">17/11/2000</div>
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016		Authorized officer <div style="text-align: center; font-weight: bold;">Beyss, E</div>

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 1,2 (partially)

In present claims 1 and 2 mentioned expressions "water dispersible compound" and "ester" are too broad and relate to an extremely large number of possible products. Support within the meaning of Article 6 PCT and/or disclosure within the meaning of Article 5 PCT is to be found, however, for only a very small proportion of the products claimed. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search over the whole of the claimed scope is impossible. Consequently, the search has been carried out for those parts of the claims which appear to be supported and disclosed, namely those parts relating to the "water dispersible compound" disclosed in claims 5-7 and to the "ester" disclosed in claims 8-24. The general idea of claims 1 and 2 has also been searched.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

INTERNATIONAL SEARCH REPORT

Information on patent family members

Intern/ 31 Application No
PCT/US 00/17431

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9505160 A	23-02-1995	AU 677705 B	01-05-1997
		AU 7499294 A	14-03-1995
		AU 7612294 A	14-03-1995
		BR 9407351 A	08-10-1996
		CA 2166468 A	23-02-1995
		DE 69417424 D	29-04-1999
		DE 69417424 T	15-07-1999
		WO 9505159 A	23-02-1995
		EP 0713385 A	29-05-1996
		ES 2129135 T	01-06-1999
		JP 2919973 B	19-07-1999
		JP 9501668 T	18-02-1997
		ZA 9406132 A	15-02-1996
		AU 696398 B	10-09-1998
		AU 7612194 A	14-03-1995
		BR 9407231 A	24-09-1996
		CA 2166469 A	23-02-1995
		WO 9505153 A	23-02-1995
		EP 0793481 A	10-09-1997
		JP 3020278 B	15-03-2000
		JP 9501665 T	18-02-1997
		ZA 9406128 A	17-07-1995
WO 9835652 A	20-08-1998	DE 19705822 A	27-08-1998
		DE 19724587 A	24-12-1998
		BR 9805982 A	31-08-1999
		EP 0897299 A	24-02-1999
US 5888951 A	30-03-1999	FR 2755369 A	07-05-1998
		DE 69700093 D	18-02-1999
		DE 69700093 T	27-05-1999
		EP 0839521 A	06-05-1998
		ES 2129277 T	01-06-1999
		JP 2986436 B	06-12-1999
		JP 10139626 A	26-05-1998
WO 9505145 A	23-02-1995	AU 677704 B	01-05-1997
		AU 7499194 A	14-03-1995
		BR 9407229 A	24-09-1996
		CA 2166532 A	23-02-1995
		DE 69408148 D	26-02-1998
		DE 69408148 T	07-05-1998
		EP 0713381 A	29-05-1996
		ES 2111323 T	01-03-1998
		JP 2944757 B	06-09-1999
		JP 9501667 T	18-02-1997
		ZA 9406131 A	15-02-1996